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### **FINAL**

Confirmation Sampling and Analysis Plan for the BX Service Station, Facility 736, Site ST-29



Patrick Air Force Base Florida

**Prepared For** 

Air Force Center for Environmental Excellence Brooks Air Force Base, Texas

and

45 CES/CEV Patrick Air Force Base, Florida

June 1998



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#### **FINAL**

# CONFIRMATION SAMPLING AND ANALYSIS PLAN FOR THE BX SERVICE STATION, FACILITY 736, SITE ST-29

# PATRICK AIR FORCE BASE FLORIDA

#### Prepared for:

Air Force Center for Environmental Excellence Brooks Air Force Base, Texas

and

45 CES/CEV Patrick Air Force Base, Florida

June 1998

Prepared by:

Parsons Engineering Science, Inc. 1700 Broadway, Suite 900 Denver, Colorado 80290

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#### ACRONYMS AND ABBREVIATIONS

45 CES/CEV 45th Civil Engineering Squadron/Environmental Flight

AFB Air Force Base

AFCEE Air Force Center for Environmental Excellence

AS Air Station

AST aboveground storage tank bgs below ground surface

BTEX benzene, toluene, ethylbenzene, and xylenes

BX Base Exchange

CAR contamination assessment report COPC chemical of potential concern

CPT cone penetrometer

CW compliance monitoring well

EP extraction procedure
ES Engineering-Science, Inc.

ESE Environmental Science & Engineering, Inc.

FAC Florida Administrative Code

FDEP Florida Department of Environmental Protection

HVW horizontal vent well

ICE internal combustion engine
LIF laser-induced fluorescence
LNAPL light nonaqueous-phase liquid

long-term monitoring LTM micrograms per kilogram  $\mu g/kg$  $\mu g/L$ micrograms per liter mg/kg milligrams per kilogram milligrams per liter mg/L **MOGAS** motor vehicle gasoline MP monitoring point **MTBE** methyl tert-butyl ether **NFA** no further action

NRMRL National Risk Management Research Laboratory

PAH polynuclear aromatic hydrocarbon PAMP Petroleum Action Management Plan Parsons ES Parsons Engineering Science, Inc.

PID photoionization detector POC point of compliance

ppmv parts per million, volume per volume

QC quality control RAP remedial action plan

SAP Sampling and Analysis Plan

SVE soil vapor extraction

TCLP toxicity characteristic leaching procedure

TPH total petroleum hydrocarbons

TRPH	total recoverable petroleum hydrocarbons
TVH	total volatile hydrocarbons
TVHA	total volatile hydrocarbon analyzer
USCS	Unified Soil Classification System
USEPA	US Environmental Protection Agency
UST	underground storage tank
VOC	volatile organic compound

#### **SECTION 1**

#### INTRODUCTION

This confirmation sampling and analysis plan (SAP) for the Base Exchange (BX) Service Station, Facility 736, Site ST-29, at Patrick Air Force Base (AFB), Florida has been prepared by Parsons Engineering Science, Inc. (Parsons ES) for submittal to the Florida Department of Environmental Protection (FDEP); the US Air Force Center for Environmental Excellence (AFCEE), Brooks AFB, Texas; and 45th Civil Engineering Squadron/Environmental Flight (45 CES/CEV), Patrick AFB, Florida. The SAP is intended to guide soil sampling at the site to document the effectiveness of remediation of petroleum-hydrocarbon-contaminated soils using *in situ* bioventing. The BX Service Station has been in service since 1954, and is the primary on-Base "gas station" used by military personnel and their families. Petroleum contamination has been identified in site soils and groundwater primarily as a result of leaks in former motor vehicle gasoline (MOGAS) underground storage tanks (USTs) and the associated piping. There is no evidence of mobile light nonaqueous-phase liquid (LNAPL, "free-phase product") at this site.

In 1992, the BX Service Station was selected as a pilot test site for the AFCEE Bioventing Initiative. The purpose of this initiative was to determine if *in situ* bioventing would be a feasible cleanup technology for source area petroleum-contaminated soils within the unsaturated zone. Under the initiative, Parsons ES (formerly Engineering-Science, Inc. [ES, 1993]) installed a pilot-scale bioventing system at the BX Service Station in March 1993, and performed initial pilot testing. Prior to extended pilot testing using air injection bioventing, soil vapor extraction (SVE) was performed for 3 months to reduce high concentrations of total volatile hydrocarbons (TVH) in vadose zone soils. Following SVE and a 96-percent reduction of TVH in extracted soil gas, the system was reconfigured for air injection bioventing and operated for an additional 9 months prior to 1-year testing. Results of the 1-year pilot test demonstrated that SVE and air injection bioventing had effectively treated petroleum-contaminated soils in the area influenced by the pilot-scale system.

In September 1994, the BX Service Station was included in the AFCEE Extended Bioventing project and funded for system expansion (Option 4), extended bioventing system operation followed by soil gas sampling and *in situ* respiration testing (Option 1), and confirmation/closure sampling (Option 2) following sufficient bioventing treatment of site soils. In November 1996, soil gas sampling and respiration testing was performed following approximately 15 months of expanded bioventing system operation, and results indicated that benzene, toluene, ethylbenzene, and xylenes (BTEX) concentrations in vadose zone soils had been significantly reduced (Parsons ES, 1997a). An additional 6 months to 1 year of continued system operation was

recommended by Parsons ES and AFCEE, prior to initiating confirmation/closure sampling activities, because of elevated TVH concentrations in soil gas at one location. To date, soils in the vicinity of the original pilot-scale system have benefited from more than 3.5 years of soil venting, and soils influenced by the expanded bioventing system have received more than 2.5 years of treatment. Based on previous sampling results and extended system operation, petroleum hydrocarbon concentrations in vadose zone soils are likely to meet revised FDEP (1997) soil target cleanup levels which appear in Chapter 62-770 of the Florida Administrative Code (FAC), as listed in Section 3 of this SAP.

The objective of the confirmation sampling is to document the effectiveness of *in situ* bioventing for the remediation of petroleum-hydrocarbon contaminated soils. The site is currently under a long-term monitoring status. The proposed confirmation sampling described in Section 4 targets unsaturated and smear zone soils in the vicinity of the current fuel dispensing island and the MOGAS USTs. It is anticipated that soil analytical results will meet FDEP (1997) risk-based criteria for NFA, and that no further soil remediation will be necessary.

This SAP consists of nine sections, including this introduction, and one appendix. Section 2 includes a site description and history, and summaries of previous investigations and remediation activities. Section 3 summarizes cleanup requirements for the BX Service Station. A detailed SAP is presented in Section 4. Analytical results will be presented in a confirmation sampling report, as described in Section 5. Section 6 lists Patrick AFB support requirements, and Section 7 presents the proposed project schedule. Air Force, regulatory, and contractor points of contact are provided in Section 8, and references for this SAP are provided in Section 9. Appendix A provides pertinent groundwater analytical data from previous sampling events.

#### **SECTION 2**

#### SITE DESCRIPTION

#### 2.1 SITE LOCATION AND PROJECT HISTORY

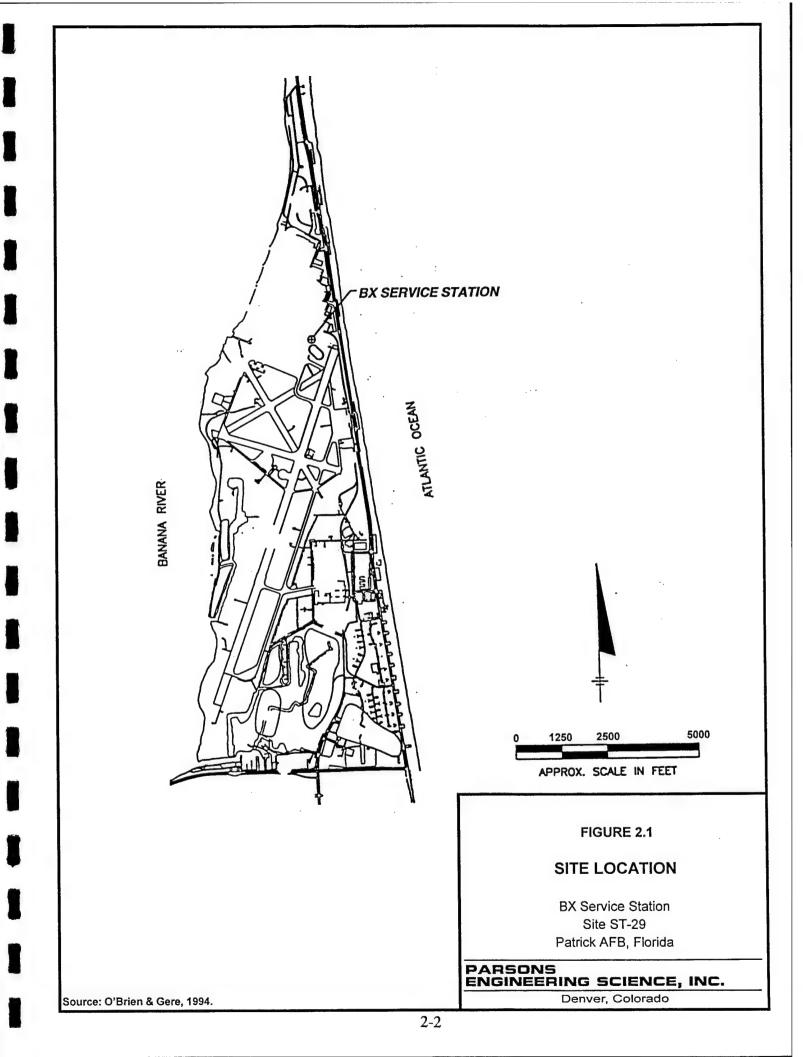
Patrick AFB lies on a long barrier island situated off Florida's east coast, in Brevard County. Patrick AFB is bounded on the east by the Atlantic Ocean and on the west by the Banana River. The City of Cocoa Beach is located immediately north of the Base, and Satellite Beach is directly south. The BX Service Station is located in the north-central section of Patrick AFB (Figure 2.1).

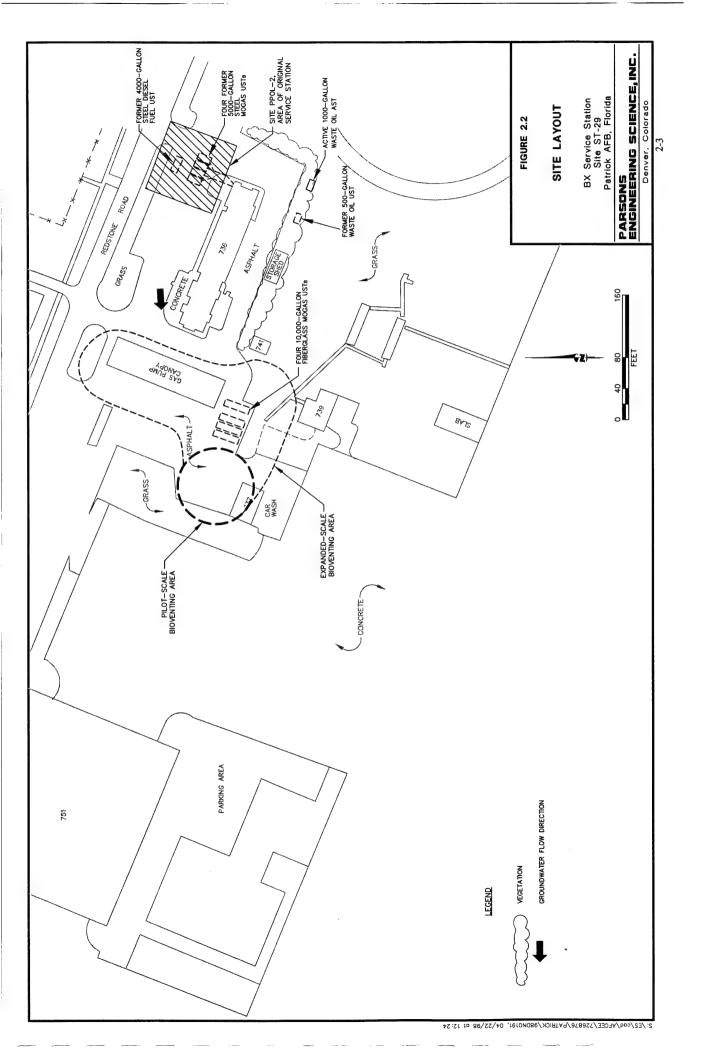
The original service station area is located to the northeast of the existing building (Building 736), as shown on Figure 2.2. The original service station (Site PPOL-2) consisted of four former 5,000-gallon steel USTs that contained MOGAS, a former 4,000-gallon steel UST that contained diesel fuel, and two former dispenser islands and the associated piping. The former dispenser islands were located above the USTs. During a major renovation in 1973, the dispenser islands were removed, the original USTs were abandoned in place, and the original service station area was paved over with asphaltic concrete. New fueling facilities consisting of dispenser islands, five 10,000-gallon fiberglass USTs for MOGAS, and associated piping were installed in their current locations west and southwest of Building 736 (Figure 2.2).

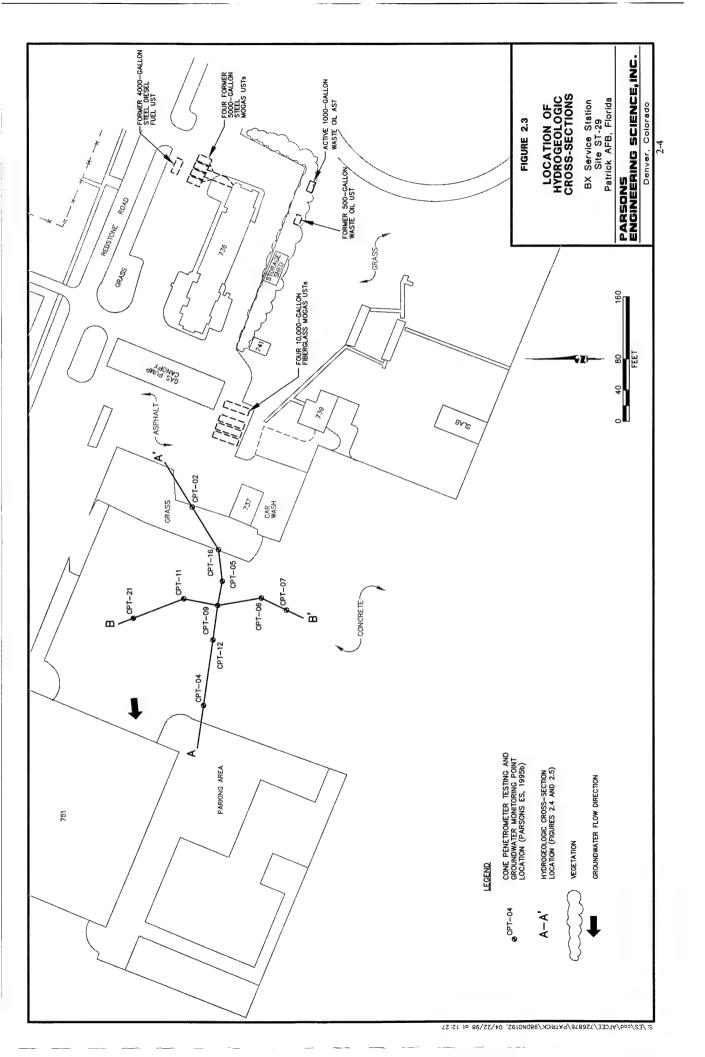
In 1985, a leak was detected in a fiberglass fuel line, and in 1986, one of the 10,000-gallon fiberglass USTs was removed due to leakage. It is estimated that approximately 700 gallons of MOGAS was released into the subsurface as a result of the fuel line and UST leaks (Environmental Science & Engineering, Inc., [ESE], 1991). In February 1992, a 500-gallon UST southeast of Building 736 was removed and replaced with a 1,000-gallon waste oil aboveground storage tank (AST). In 1993, the five abandoned USTs located northeast of Building 736 were excavated and removed along with some excessively contaminated soils (O'Brien & Gere Engineers, Inc. [OB&G], 1995).

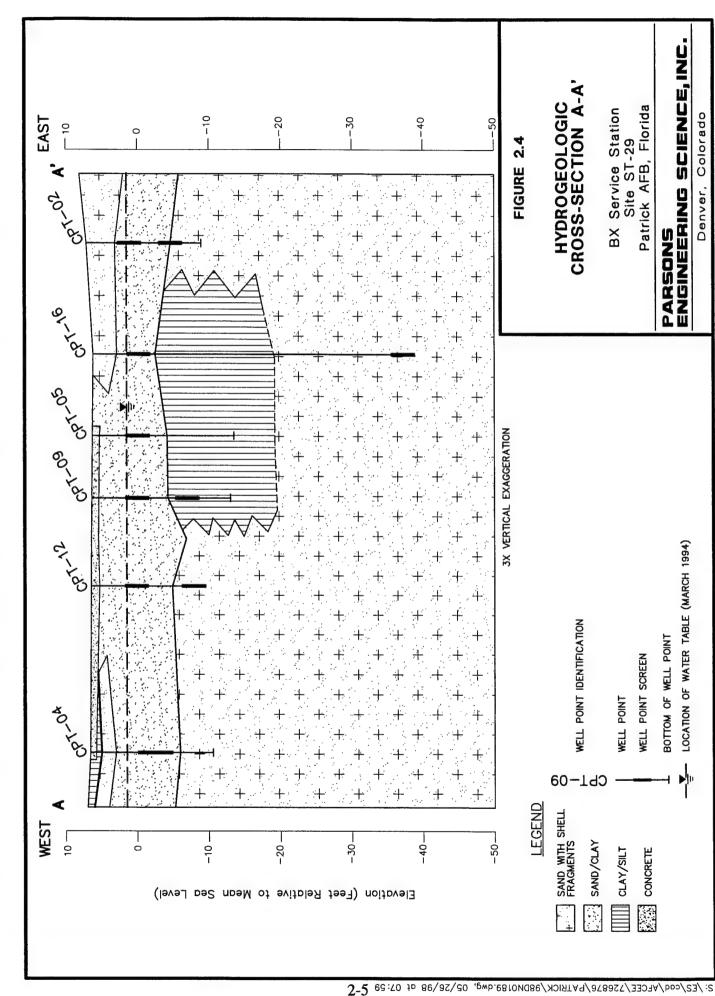
#### 2.2 SITE GEOLOGY AND HYDROGEOLOGY

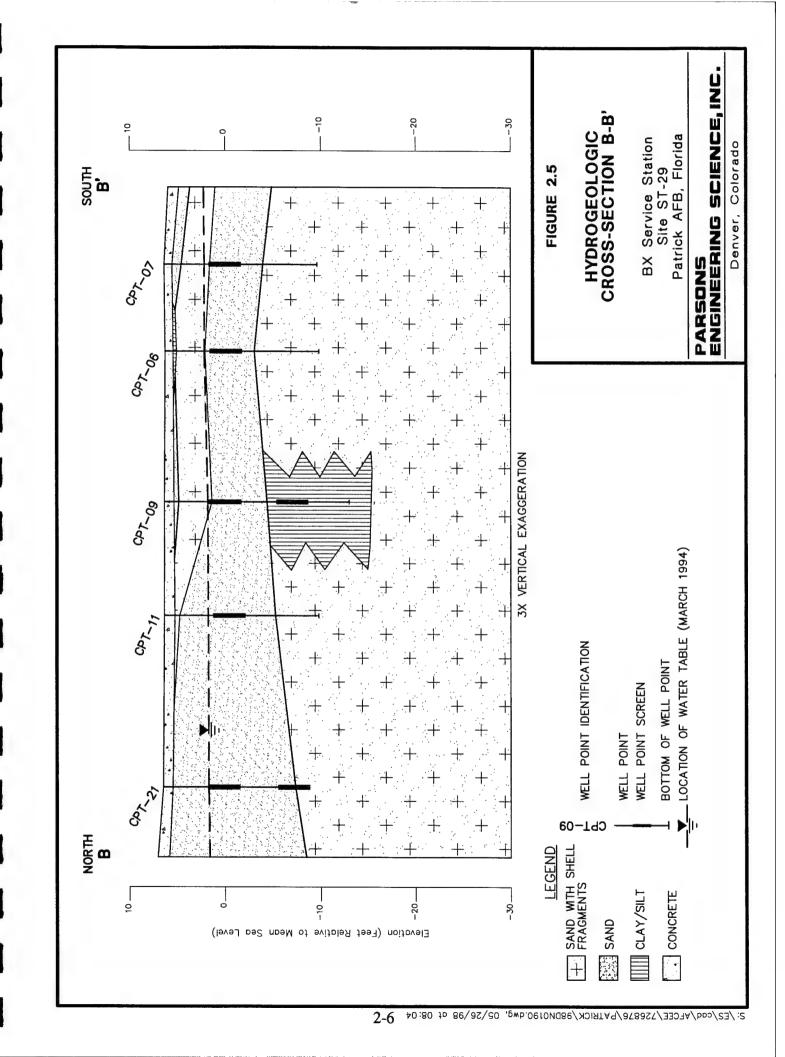
The BX Service Station is characterized by a level ground surface covered primarily with asphalt and concrete pavement. Soils at this site consist of predominantly unconsolidated, poorly to moderately well-sorted, fine- to coarse-grained quartz sand with up to 40-percent shell fragments. This marine sand deposit extends to approximately 25 feet below ground surface (bgs). Figure 2.3 shows the locations of two hydrogeologic cross-sections, which are shown on Figures 2.4 and 2.5.











Shallow groundwater at the site is unconfined and typically encountered in the sand at approximately 4 to 6 feet bgs. Historical groundwater elevation data indicate that groundwater flow at the site is generally to the west (ESE, 1991; Parsons ES, 1995b). The horizontal hydraulic gradient at the site ranges from 0.00096 to 0.003 foot per foot (ESE, 1991; OB&G, 1992; Parsons ES, 1995b). The advective groundwater velocity is estimated to be 160 feet per year (Parsons ES, 1995b). Groundwater at Patrick AFB has been designated as Class G-II under State of Florida classifications (ESE, 1991). Class G-II groundwater is acceptable for potable use, with a total dissolved solids content less than 10,000 milligrams per liter [mg/L]). However, shallow groundwater at the site and groundwater at Patrick AFB is not used for potable drinking water. The Base gets its potable water supply from the City of Cocoa (OB&G, 1995).

#### 2.3 PREVIOUS INVESTIGATIONS

#### 2.3.1 1990 Phase II, Stage 2 Investigation by ESE

As a result of the leaking pipe discovered in 1985, and the removal of one 10,000-gallon fiberglass MOGAS UST in 1986, a Phase II, Stage 2 investigation was conducted by ESE (1991) in the western and northwestern areas of the BX Service Station near the active USTs and dispenser island (Figure 2.2). During this investigation, soil samples were collected from eight boreholes (SB-1 through SB-8); five shallow monitoring wells (PPOL2-1 through PPOL 2-5) and one deep monitoring well (PPOL2-6) were installed; and two rounds of groundwater samples were collected. Soil sample results from this investigation are shown in Table 2.1, and groundwater sample results are presented in Appendix A. Soil and groundwater sampling locations are shown on Figure 2.6.

Soil samples were analyzed for total recoverable petroleum hydrocarbons (TRPH), lead, volatile organic compounds (VOCs), and leachable extraction-procedure (EP) toxic metals. Elevated concentrations of TRPH, toluene, ethylbenzene, and xylenes were detected in samples collected from Boreholes SB-2 (adjacent to Building 737), SB-3 (located south of the four active MOGAS USTs), and SB-8 (adjacent to the active dispensing island). The most significant hydrocarbon contamination was evident at SB-2, where TRPH, ethylbenzene, toluene, and xylenes were detected at concentrations of 386 milligrams per kilogram (mg/kg), 100 mg/kg, 38 mg/kg, and 822 mg/kg, respectively.

Groundwater samples were analyzed for TRPH, total and dissolved lead, purgeable halocarbons, and purgeable aromatic compounds. During Round 2 sampling at monitoring well PPOL2-1, which is located directly west of the existing dispenser islands, the fuel additive methyl tert-butyl ether (MTBE) was detected at 151 micrograms per liter ( $\mu$ g/L). At monitoring well PPOL2-3, located southwest of the original service station area, total lead was detected at 0.0221 mg/L, and dissolved lead was detected at 0.0029 mg/L.

#### 2.3.2 Original Service Station Investigation and Remediation

A site characterization of the original service station area (Site PPOL-2) northeast of Building 736 (Figure 2.6) was performed by OB&G (1994 and 1995) from April 1993

TABLE 2.1
ESE AND OB&G SOIL ANALYTICAL RESULTS (1990 AND 1993)
BX SERVICE STATION
SITE ST-29
PATRICK AFB, FLORIDA

								SB-15	(4-6)		1	1	;	1	1	752		;	9	28	62	:	1
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San	SB-3	(NA)	2.16	37.3	24	22	155	SB-10	(2-4)		1	1	!	ŀ	1	39.7		0.20J	1	;	ı	0.12J	0.14J
	SB-2	(NA)	5.44	386	100	38	822	6-	(4-6)		;	:	1	0.004	;	130		1	;	1	1	1	1
	SB-1	$(NA^{c})$	5.77	58.3	;	1	22.3	SB-9	(2-4)		1	:	:	;	:	62.3		1	1	:	1	;	ı
Analyte (mg/kg) <sup>2/</sup>		ESE (1991) <sup>b/</sup>	Lead	TRPH	Ethylbenzene	Toluene	Xylenes	•	O'Brien & Gere (1994) <sup>e/</sup>	Volatiles <sup>g/</sup>	Acetone	Benzene	Ethylbenzene	Methylene Chloride	Toluene	TRPH <sup>j/</sup>	Semiyolatiles <sup>W</sup>	Fluoranthene	Naphthalene	1-Methylnaphthalene	2-Methylnaphthalene	Phenanthrene	Pyrene

" mg/kg = milligrams per kilogram.

<sup>b/</sup> Soil sample results from 1990 Phase II, Stage 2 investigation.

o' Sample depths not available.

d"--" = not detected, detection limit not available.

of Soil sample results from 1993 Petroleum Contamination Assessment, only those analytes that were detected are shown. Soil samples were collected prior to tank and soil excavation.

 $^{\prime\prime}$  D = sample replicate.

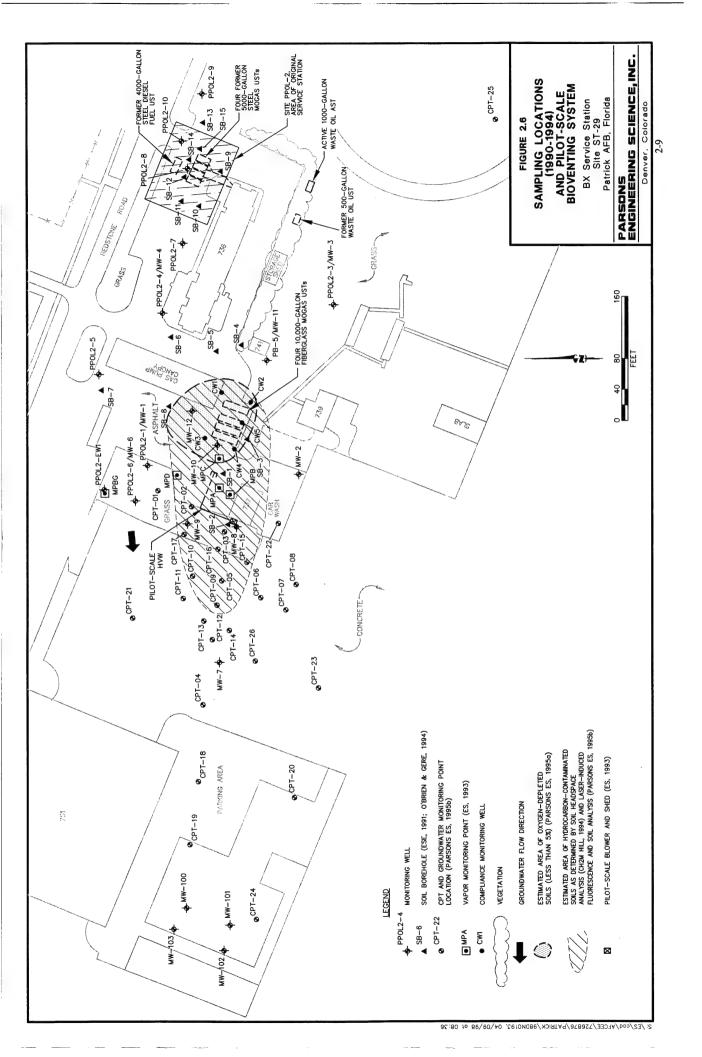
B Analyzed by USEPA Method SW8240.

 $\mathbf{b}^{\mathbf{b}}$   $\mathbf{J}$  = laboratory estimated concentration.

 $^{i\prime}$  Blank result means sample not analyzed for this parameter.

<sup>j</sup>/ Analyzed by USEPA Method SW9073.

<sup>▶</sup> Analyzed by USEPA Method SW8270.



through July 1994 to evaluate the horizontal and vertical extent of soil contamination, and to assess the potential effects of past releases of MOGAS and/or diesel fuel on shallow groundwater. Results of this study were provided in a Contamination Assessment Report (CAR) for Site PPOL-2 (OB&G, 1995). Soil results are summarized on Table 2.1. Based on soil (Boreholes SB-9 through SB-15) and groundwater (wells PPOL2-4, PPOL2-5, PPOL2-7, PPOL2-8, and PPOL2-9) analytical results from this investigation, petroleum contamination was determined to be most significant in the area of the abandoned USTs and did not extend significantly downgradient. Maximum concentrations of naphthalene (60 mg/kg), 1-methylnaphthalene (28 mg/kg), and 2-methylnaphthalene (62 mg/kg) in soil were detected in the 4- to 6-foot sample interval at SB-15. Elevated concentrations of these polynuclear aromatic hydrocarbons (PAHs) were also detected in the 4- to 6-foot sample interval at SB-12.

Following site investigation activities, the four abandoned MOGAS USTs and one abandoned diesel UST, and approximately 200 tons of excessively contaminated soils were removed between November 30 and December 3, 1993, to eliminate the source of soil and groundwater contamination at the original service station area. Results of the UST closure were documented in a facility engineering report and included as an appendix to the CAR (OB&G, 1995). In January 1996, FDEP (1996) approved an NFA proposal and issued a Site Rehabilitation Completion Order for the original service station area (Site PPOL-2).

#### 2.3.3 March 1994 CH2M Hill Soil Headspace Analysis

In March 1994, a soil headspace analyses investigation was conducted by CH2M Hill (1994) in the vicinity of the active pump islands and USTs west and southwest of Building 736, respectively (Figure 2.6). The soil headspace sampling was performed at 25 locations in accordance with FDEP soil screening guidelines for the headspace analysis method prescribed in FAC, Chapter 17-770.200 (2). The soil vapor VOC headspace readings ranged from 0 to 3,244 parts per million, volume per volume (ppmv) and indicated that soil contamination was most significant immediately adjacent to the pump islands and UST locations.

#### 2.3.4 1994 and 1995 Natural Attenuation Study

As part of a nationwide, multi-site demonstration, Parsons ES (1995b), in conjunction with researchers from the US Environmental Protection Agency (USEPA) National Risk Management Research Laboratory (NRMRL; formerly Robert S. Kerr Environmental Research Laboratory), was retained by AFCEE to conduct site characterization and groundwater modeling at the BX Service Station (Site ST-29) to evaluate the effectiveness of natural attenuation of dissolved fuel contaminants. To support the study, site characterization data, including soil and groundwater sampling and aquifer testing, were collected in March 1994 (by Parsons ES and NRMRL), March 1995 (NRMRL), and May 1995 (CH2M Hill). Cone penetrometer testing (CPT) was conducted to collect stratigraphic information and soil samples in the vicinity of the active pump islands and USTs and downgradient from the site. Groundwater samples were collected at monitoring points installed in CPT boreholes,

and at previously installed monitoring wells. Soil and groundwater sampling locations are shown on Figure 2.6.

Soil samples collected in March 1994 were analyzed for BTEX, total petroleum hydrocarbons (TPH), trimethylbenzene isomers, and total organic carbon. Soil BTEX and TPH data are included in Table 2.2. The highest concentrations of total BTEX and TPH observed during this investigation were from samples collected near the water table (e.g., 5.5 to 6.5 feet bgs) downgradient from the active USTs and pump islands. A total BTEX concentration of 1,236 mg/kg and a TPH concentration of 17,100 mg/kg were detected in the sample collected 6 feet bgs at CPT-02. At CPT-03, a TPH concentration of 11,700 mg/kg was detected in the sample collected 5.5 feet bgs, and a total BTEX concentration of 975 mg/kg was detected in the 6.5-foot bgs soil sample. Total BTEX concentrations are generally an order of magnitude lower in samples collected 1 to 2 feet above or below these capillary fringe samples. During this investigation, the downgradient extent of smear zone hydrocarbon contamination was estimated to be approximately 220 feet downgradient from the source area (Figure 2.6), where total BTEX and TPH concentrations of 0.0989 mg/kg and 2,740 mg/kg, respectively, were detected in the 5-foot-bgs sample from CPT-09.

Groundwater samples collected in March 1994 (by Parsons ES and NRMRL), in March 1995 (by NRMRL), and in May 1995 (by CH2M Hill) confirmed the presence of fuel-hydrocarbon contamination in the shallow saturated zone in the vicinity of the BX Service Station and extending downgradient to an area just south of Building 751 (Figure 2.6). Dissolved BTEX contamination has migrated west of the active USTs and pump islands in the direction of groundwater flow; however, data collected between March 1994 and March/May 1995 indicate that natural chemical attenuation processes have been effective in preventing further downgradient migration of the dissolved BTEX plume. Groundwater contaminant and geochemical data indicate that biodegradation is reducing dissolved contaminant mass, and the BTEX plume shrank in both the longitudinal and lateral directions between March 1994 and March/May 1995 (Parsons ES, 1995b). Tables and figures summarizing groundwater contaminant data and distribution based on these sampling events are provided in Appendix A.

The results of the study suggest that natural attenuation of dissolved BTEX compounds is occurring at sufficient rates to reduce concentrations below regulatory guidelines before potential downgradient receptors could be adversely affected. The study recommended the use of long-term monitoring (LTM) wells and point-of-compliance (POC) monitoring wells to monitor the long-term migration and degradation of the dissolved BTEX plume (Parsons ES, 1995b). The study also recommended continued bioventing until source area contamination in site soils was sufficiently reduced.

#### 2.4 PILOT-SCALE AND EXPANDED-SCALE BIOVENTING

As part of the AFCEE Bioventing Initiative, ES (1993) installed a pilot-scale bioventing system at the BX Service Station in March 1993 to assess the potential for air injection bioventing to remediate the hydrocarbon contamination identified in vadose zone soils. The primary objectives of the pilot test were: 1) to assess the potential for

# TABLE 2.2 PARSONS ES SOIL ANALYTICAL RESULTS (1993-1995) BX SERVICE STATION

#### SITE ST-29 PATRICK AFB, FLORIDA

				Analyte <sup>a/</sup>		
Sampling Event/	Depth	TPH	Benzene	Toluene	Ethylbenzene	Xylenes
Sample Location	(ft bgs) <sup>b/</sup>	(mg/kg) <sup>c/</sup>	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)
March 1994 Natural						
Attenuation Study <sup>d/</sup>						
CPT-02-A17	4	ND <sup>e/</sup>	<0.02 <sup>f</sup>	< 0.02	0.0345	0.1085
CPT-02-A18	5	140	0.197	0.0605	0.514	3.119
CPT-02-A19	6	17,100	6.99	8.08	191	1,030
CPT-02-A20	6.45	1,660	0.68	0.345	16.1	84.1
CPT-02-A21	7	289	0.339	0.106	2.26	12.42
CPT-03-A22	3	20.9	0.164	0.0142	0.21	1.019
CPT-03-A23	4.5	0.2	ND	< 0.02	0.00838	0.0418
CPT-03-A24	5.5	11,700	< 0.02	0.462	< 0.02	171.3
CPT-03-A25	6.5	9,300	2.33	29.5	120	823
CPT-03-A26	7	287	0.366	0.532	2.02	15.46
CPT-05-A1	3.5	ND	< 0.02	< 0.02	< 0.02	< 0.02
CPT-05-A2	4.5	ND	< 0.02	< 0.02	< 0.02	< 0.02
CPT-05-A3	5	ND	< 0.02	< 0.02	< 0.02	0.00725
CPT-05-A4	5.5	ND	< 0.02	< 0.02	< 0.02	< 0.02
CPT-09-A4	2.5	2	< 0.02	< 0.02	ND	< 0.02
CPT-09-A5	4.5	47.1	< 0.02	< 0.02	< 0.02	< 0.02
CPT-09-A6	5	2,740	< 0.02	0.034	< 0.02	0.0649
CPT-09-A7	6	15.2	< 0.02	< 0.02	< 0.02	< 0.02
CPT-13	5.5	ND	< 0.02	< 0.02	< 0.02	< 0.02
CPT-15-A8	3	ND	< 0.02	< 0.02	ND	< 0.02
CPT-15-A9	4	0.01	< 0.02	< 0.02	ND	< 0.02
CPT-15-A10	5	526	< 0.02	0.012	0.00978	0.0226
CPT-15-A11	5.5	ND	ND	ND	< 0.02	< 0.02
CPT-16-A1	4.5	0.04	0.00752	0.00761	< 0.02	< 0.02
CPT-16-A2	6	ND	ND	< 0.02	ND	< 0.02
CPT-17-A12	2	ND	ND	< 0.02	< 0.02	< 0.02
CPT-17-A13	3	ND	ND	< 0.02	ND	< 0.02
CPT-17-A14	4	ND	ND	< 0.02	< 0.02	0.00779
CPT-17-A15	5	ND	ND	< 0.02	ND	< 0.02

## TABLE 2.2 (Continued) PARSONS ES SOIL ANALYTICAL RESULTS (1993-1995)

#### **BX SERVICE STATION**

#### SITE ST-29

#### PATRICK AFB, FLORIDA

				Analyte <sup>a/</sup>		
Sampling Event/	Depth	TRPH	Benzene	Toluene	Ethylbenzene	Xylenes
Sample Location	(ft bgs) <sup>b/</sup>	(mg/kg) <sup>c/</sup>	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)
Pilot-Scale Bioventing March 1993 <sup>g/</sup>						
HVW-2	4.5	2,730	< 14 <sup>h/</sup>	54	260	2,600
MPA	3.5	11	< 6.2	23	320	140
MPC	3.5	60	< 0.31	< 0.36	< 0.26	5.7
December 1994 <sup>i</sup>						
HVW-2	4.5	81.9	< 0.05	< 0.05	< 0.05	< 0.099
MPA	3.5	50.8	< 0.049	< 0.049	< 0.049	< 0.098
MPC	3.5	57.6	0.13	0.15	0.16	0.49
Expanded-Scale Bioventi July 1995 <sup>j/</sup>	ing					
MPE	3.5	743	0.075	< 0.05	20	160
MPF	3.5	767	0.47	0.57	7.4	9.0
CW2	3.5	13.3	< 0.05	< 0.05	< 0.05	< 0.13

<sup>&</sup>lt;sup>a'</sup> For natural attenuation study, total petroleum hydrocarbons (TPH) quantified with a JP-4 jet fuel standard. For pilot-scale and expanded scale bioventing, total recoverable petroleum hydrocarbons (TRPH) analyzed by USEPA Method 418.1; benzene, toluene, ethylbenzene, and total xylenes analyzed by USEPA Method SW8020.

b' ft bgs = feet below ground surface.

e' mg/kg = milligrams per kilogram.

d Parsons ES, 1995a.

<sup>&</sup>lt;sup>e'</sup> ND = not detected and detection limit not available.

g < = below limit of quantitation shown for natural attenuation study results.

g/ ES, 1993. Soil samples collected prior to pilot-scale bioventing system startup.

<sup>&</sup>lt;sup>b/</sup> <= analyte concentration in sample was less than the method detection limit shown for bioventing results.

<sup>&</sup>lt;sup>1</sup> Soil samples collected following approximately 3 months of soil vapor extraction and 9 months of pilot-scale bioventing system operation.

<sup>&</sup>lt;sup>j</sup> Soil samples collected during full-scale bioventing system installation.

supplying oxygen throughout the contaminated soil profile; 2) to determine the rate at which indigenous microorganisms would degrade petroleum hydrocarbons when stimulated by oxygen-rich soil gas at this site; and 3) to evaluate the potential for sustaining these rates of biodegradation until hydrocarbon contamination is remediated below regulatory approved standards (ES, 1993).

The pilot test location and system configuration were based on results of a soil gas survey performed by ES in January 1993 in the vicinity of SB-2. The soil gas survey indicated TVH concentrations exceeding 10,000 ppmv and depleted oxygen conditions in soils immediately north of the car wash, Building 737 (ES, 1993). The pilot-scale bioventing system consisted of one horizontal vent well (HVW-2), five vapor monitoring points (MPA, MPB, MPC, MPD, and MPBG), a 1-horsepower regenerative blower, blower piping and gauges, and a weatherproof blower shed (see Figure 2.6). The HVW was installed at the site at a depth of approximately 4 feet bgs because of the relatively shallow water table. At each of the MPs, a single screened interval was placed from 3.0 to 3.5 feet bgs. During installation and testing of the pilot-scale system, soil and soil gas sampling, and respiration and soil-to-air permeability testing were performed. Based on results of the oxygen influence and air permeability testing, the long-term radius of oxygen influence around the HVW was expected to exceed 37 feet. A detailed description of the pilot-scale bioventing system design and initial testing results are provided in the Draft Interim Pilot Test Results Report (ES, 1993).

Prior to extended operation of the pilot-scale air injection bioventing system, SVE was performed at the BX Service Station site to reduce the potential for uncontrolled VOC vapor migration and discharge to the atmosphere. Soil gas samples collected during the January 1993 soil gas survey and initial soil gas samples collected following pilot-scale bioventing system installation indicated significant concentrations of TVH in soil gas at the site, and the need for a short period of SVE prior to air injection bioventing. The SVE system utilized a modified internal combustion engine (ICE) for the removal and destruction of hydrocarbon vapors extracted from fuel-contaminated soils at the site (AFCEE, 1994). The SVE system was operated from October 18, 1993 through January 14, 1994, when the blower was reconfigured for air injection Startup of the bioventing system immediately followed SVE system bioventing. shutdown and continued until November 1994. One-year respiration testing and soil gas sampling was performed in December 1994, following 1 month of system shutdown to allow equilibrium conditions to develop in site soils, for comparison to initial results. Following the 1-year testing event, the system was started and reoptimized for continuous air injection.

In December 1994, during the collection of 1-year data for the pilot-scale system, Parsons ES performed a soil gas survey in the vicinity of the active pump islands and USTs to supplement the March 1994 CH2M Hill investigation (Section 2.3.3). The results from this survey indicated that the extent of remaining unsaturated soil contamination was localized around the active pump islands and USTs west and southwest of Building 736, and that soil gas oxygen levels in this area were depleted (less than 5 percent). The estimated area of oxygen depletion based on this survey is shown on Figure 2.6.

Based on favorable 1-year testing results for the pilot-scale bioventing system, and the presence of contaminated soil outside the effective treatment area of HVW-2, the BX Service Station site was added to the AFCEE Extended Bioventing Project (Contract No. F41624-92-D-8036, Order 17, awarded by AFCEE to Parsons ES on 30 September 1994). Under the extended bioventing project, funding was allocated for bioventing system expansion at the site (Option 4); for an additional year of system operation and testing (Option 1); and for confirmation/closure soil sampling (Option 2), following sufficient remediation of petroleum-hydrocarbon-contaminated soils.

In preparation for bioventing system expansion, Parsons ES (1995a) developed an Initial Remedial Action Plan and performed a supplemental soil headspace survey in May 1995. The survey further delineated the soils to be remediated through bioventing system expansion. Following Patrick AFB, AFCEE, and FDEP approval of the Initial Remedial Action Plan, the expanded full-scale bioventing system was installed in July 1995 in the vicinity of the active pump islands and USTs. The expanded-scale system consists of two additional HVWs (HVW-1 and HVW-3), five additional MPs (MPE, MPF, MPG, MPH, and MPI), and a new blower system. A layout of the expanded-scale bioventing system is presented on Figure 2.7.

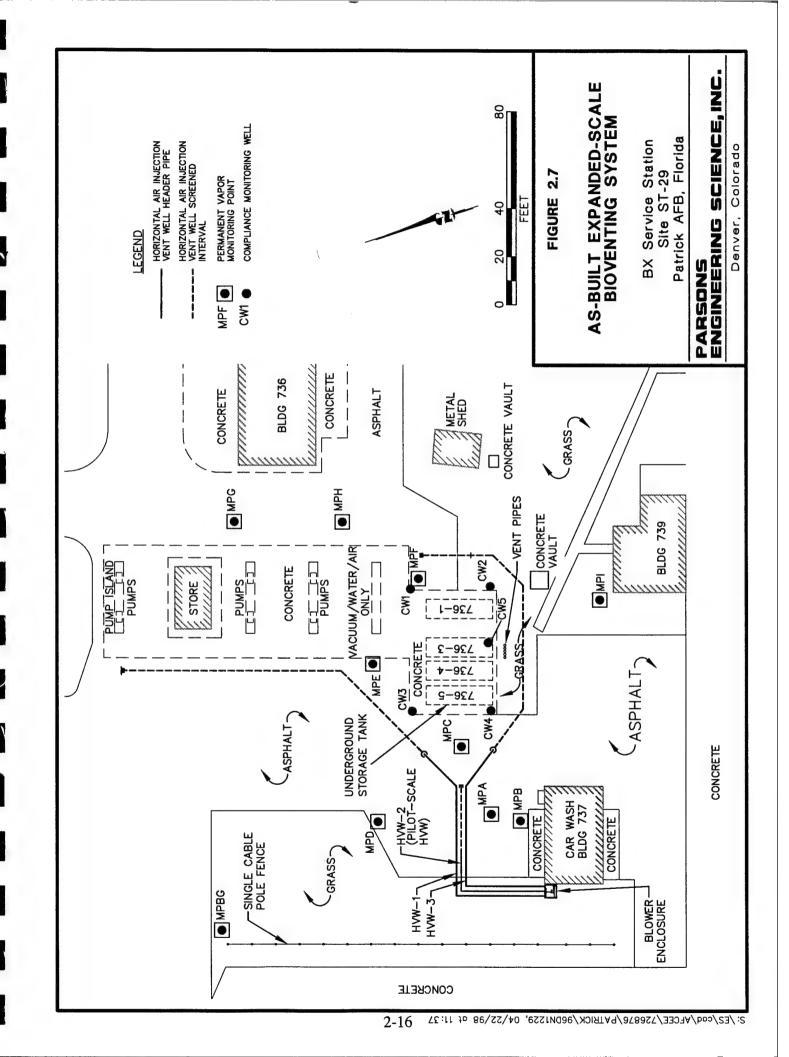
Expanded-scale system operation began on July 12, 1995. Option 1 soil gas sampling and respiration testing was performed in early November 1996, following 1 month of system shutdown to allow equilibrium conditions to develop in site soils and to allow comparison to initial and 1-year results. Results of the Option 1 testing event were presented in a Parsons ES (1997a) letter report to AFCEE and Patrick AFB. The expanded-scale bioventing system was restarted following Option 1 testing. In March 1997, Parsons ES revisited the site to perform a systems maintenance check. Site measurements demonstrated that the entire area designated for treatment was receiving oxygen as a result of expanded-scale bioventing system operation (Parsons ES, 1997b).

#### 2.4.1 Soil Sampling Results

Soil sampling by Parsons ES was conducted at the BX Service Station site during installation of the pilot-scale bioventing system (March 1993), after 12 months of SVE/pilot-scale air injection bioventing (December 1994), and during installation of the full-scale bioventing system (July 1995). A summary of the pilot-scale and expanded-scale soil analytical results is provided in Table 2.2.

The TRPH concentrations in soils sampled from the immediate vicinity of the pilot-scale horizontal vent well decreased 1 to 2 orders of magnitude between March 1993 and December 1994. TRPH concentrations at MPA and MPC showed little contamination prior to SVE and pilot-scale bioventing, and remained low at the 1-year sampling event. BTEX concentrations in soils at HVW-2 and MPA decreased 3 to 4 orders of magnitude to nondetect levels as a result of SVE and air injection bioventing. Lesser decreases in BTEX soil concentrations were evident at MPC.

Soil samples also were collected from MPE, MPF, and compliance monitoring well CW2 (Figure 2.7) in July 1995 during bioventing system expansion. TRPH and BTEX concentrations in soils collected from MPE and MPF indicated these areas were moderately impacted by fuel hydrocarbon contamination. Significant contamination



was not evident at CW2. Although no soil sampling was performed by Parsons ES following 1 year of full-scale system operation, based on measured reductions in soil gas TVH and BTEX concentrations and decreased *in situ* respiration rates, significant reductions in soil TRPH and BTEX concentrations appear to have occurred as the result of expanded-scale bioventing system operation.

#### 2.4.2 Soil Gas Sampling Results

Soil gas sampling was performed in March 1993 prior to SVE/bioventing system startup (in October 1993), in December 1994 following 3 months of SVE and 9 months of air injection bioventing system operation, in July 1995 prior to expanded-scale bioventing system startup, and in November 1996 following 15 months of expanded-scale bioventing system operation. Soil gas samples were collected from each MP and field-screened to assess soil gas concentrations of oxygen, carbon dioxide, and TVH. Results presented in Table 2.3 indicate that static oxygen concentrations in soil gas have generally increased with continued bioventing at the site. During the November 1996 sampling event, soils at MPF had not experienced a measurable increase in soil gas oxygen concentration, which suggested ongoing aerobic activity associated with relatively greater residual hydrocarbon contamination (Table 2.2).

Initial, 1-year, and 3-year soil gas samples also were collected at selected locations for laboratory analysis (Table 2.3). For all three sampling events, samples were sent to the Air Toxics, Ltd. laboratory in Folsom, California and analyzed for TVH and BTEX using USEPA Method TO-3. Comparison of 1-year and 3-year soil gas results demonstrated substantial reductions in TVH and BTEX concentrations compared to initial values. With the exception of total xylenes at MPE, BTEX concentrations had been reduced to less than 1.0 ppmv at all locations sampled during the November 1996 sampling event. During the initial 3 years of soil venting system operation, TVH concentrations were reduced from between 38,000 and 100,000 ppmv to less than 1.0 ppmv in the pilot-scale area (MPB and MPC). Soil gas results from MPE and MPF, installed as part of the expanded-scale bioventing system, showed TVH concentrations were reduced between 1 and 3 orders of magnitude following the first year of expanded-scale system operation.

With the exception of results from MPF, located on the east side of the active fiberglass USTs (Figure 2.7), field and analytical soil gas results from the November 1996 sampling event strongly suggested nearly complete remediation of hydrocarbon contaminants in soils at the BX Service Station site. Results of the March 1997 maintenance check demonstrated that MPF was receiving adequate oxygen (20.3 percent) for continued aerobic biodegradation of petroleum hydrocarbon contaminants in this source area (Parsons ES, 1997b).

#### 2.4.3 Respiration Test Results

Respiration and fuel biodegradation rates for the first 3 years of soil venting are shown in Table 2.4. During each testing period, observed rates of oxygen utilization (in situ respiration) were used to estimate aerobic fuel biodegradation rates in site soils. Results of the November 1996 Option 1 testing event demonstrated that in situ respiration and fuel biodegradation rates have generally decreased with continued

SOIL GAS FIELD AND ANALYTICAL RESULTS SITE ST-29 PATRICK AFB, FLORIDA BX SERVICE STATION TABLE 2.3

			Field Screening Data	ata		¥	Analytical Data		
:	:		Carbon	Arrest		l	F	40.40	V. Image
Sampling Location"	Sampling Event	Oxygen (percent)	(percent)	(ppmv)	(hudd)	(ppmv)	(bbmv)	(ppmv)	(bpmv)
PILOT-SCALE I	PILOT-SCALE BIOVENTING SYSTEM AREA	EA							
HVW-4"	Initial (3/93)	0.0	15.0	> 20,000 <sup>0</sup>	47,000	<2.20	15	41	200
	1-Year (12/94)	<b>&gt;</b>	1	ı	!	i	ł	1	1
	3-Year (11/96)	I	i	I	•	I	;	1	1
MPA-3.5	Initial (3/93)	0.0	14.3	> 20,000	100,000	< 5.3	110	46	310
	1-Year (12/94)	11.8	5.0	230	6.9	0.033	0.003	0.008	0.46
	1.5-Year (7/95)	11.0	4.	\$0	1	I	-	i	1
	3-Year (11/96)	10.7	5.1	156	1	I	!	1	1
MPB-3.5	Initial (3/93)	0.0	13.6	> 20,000	1	i	ı	ı	I
	I-Year (12/94)	3.5	8.6	260	i	ı	***	i	1
	1.5-Year (7/95)	4.2	7.8	20	!	i	I	I	!
	3-Year (11/96)	7.5	9.9	340	0.17	< 0.002	< 0.002	< 0.002	< 0.002
MPC-3.5	Initial (3/93)	0.0	15.2	> 20,000	38,000	<2.1	8.3	12	83
	1-Year (12/94)	14.2	4.2	200	1.7	0.02	< 0.002	< 0.002	< 0.002
	1.5-Year (7/95)	11.3	0.9	40	į	ı	1	1	i
	3-Year (11/96)	12.5	5.0	164	0.13	< 0.002	< 0.002	< 0.002	< 0.002
MPD-3.5	Initial (3/93)	15.9	2.8	260	1	I	ı		1
	1-Year (12/94)	1	1	1	1	1	1	i	ł
	1.5-Year (7/95)	19.1	1.1	70	1	!	I	ŧ	•
	3-Year (11/96)	19.0	1.7	92	I	ı	I	I	I
FULL-SCALE B	FULL-SCALE BIOVENTING SYSTEM AREA	¥							
MPE-3.5	Initial (7/95) 1-Year (11/96)	0.0 8.2	18.0	961 961	27,140 9.9	88 < 0.005	130 0.11M"	240	1,200
MPF-3.5	Initial (7/95)	0.0	17.5	>20.000	27.500	86	19	300	210
	1-Year (11/96)	0.0	10.4	1,020	2,200	0.16	0.53	0.39	0.65
MPG-3.5	Initial (7/95)	15.8	4.0	10	I	i	•	•	I
	1-Year (11/96)	18.9	9:1	80	I	I	1	I	I
MPH-3.5	Initial (7/95) 1-Year (11/96)	9.8	8.0	20 88	101	< 0.002	< 0.002	< 0.002	< 0.002
MPI-3.5	fuitial (7/95)	19.5	89	10	ı	i	I	ì	ı
	1-Year (11/96)	20.0	0.7	36	ł	!	1	!	I
CW2-4"	Initial (7/95)	1	I	ı	2.29	< 0.002	0.070	< 0.002	< 0.002
	1-Year (11/96)	I	i	I	i	Ī	-	1	-

<sup>&</sup>quot;Sampling location identifies the sampled monitoring point and depth in feet below ground surface.

W TVH = total volatile hydrocarbons.

of ppmv = parts per million, volume per volume.

<sup>&</sup>lt;sup>4</sup> Soil gas analyses performed using USEPA Method TO-3.

 $<sup>^{\</sup>omega}$  The pilot-scale horizontal vent well is designated HVW-2 on Figure 2.7.  $^{\rho}$  > = measurement exceeded maximum reading for GasTech® Trace-Techtor Hydrocarbon Analyzer.

V < = analyte concentration was less than the method detection limit shown.

M= laboratory reported value may be biased due to apparent matrix interferences.

F Compliance monitoring well 2 (CW2) is screened from approximately 3 to 13 feet below ground surface. The middle of the effective screened interval for soil gas sampling was approximately 4 feet below ground surface during the 7/95 sampling event.

RESPIRATION AND FUEL BIODEGRADATION RATES BX SERVICE STATION TABLE 2.4

PA SENTICE STRAIGHT	SITE ST-29	PATRICK AFB, FLORIDA	

	Initial (N	Initial (March 1993)	6-Month (May 1994) <sup>a/</sup>	1ay 1994) <sup>a/</sup>	1-Year (Dec. 1994)	ec. 1994)	3-Year (Nov. 1996)	ov. 1996)
•	Respiration	Degradation	Respiration	Degradation	Respiration	Degradation	Respiration	Degradation
Testing	Rate	Rate	Rate	Rate	Rate	Rate	Rate	Rate
Location-Depth	(% O <sub>2</sub> /hour)	(mg/kg/year)"	(% O <sub>2</sub> /hour)	(mg/kg/year)	(% O <sub>2</sub> /hour)	(mg/kg/year)	(% O <sub>2</sub> /hour)	(mg/kg/year)
							•	8
MPA-3.5	0.17	940	0.19	970	0.029	130	Z	NC
MPB-3.5	0.15	840	0.27	1,400	0.035	150	0.084	380
MPC-3.5	0.16	970	0.16	850	0.020	92	MN	NC
MPE-3.5	/B	1	1		-	•	990.0	290
MPF-3.5			*	-	ļ	1	0.12	540

<sup>&</sup>quot;Initial bioventing pilot test occurred in March 1993, but SVE/air injection bioventing system operation did not begin until October 18, 1993.

 $<sup>^{\</sup>omega}$  Milligrams of hydrocarbons per kilogram of soil per year.  $^{\omega}$  6-month degradation rates based on average soil moisture results for initial and 1-year soil sampling events.

<sup>&</sup>lt;sup>o</sup> 3-year degradation rates based on average soil moisture result from December 1994 soil sampling event.

<sup>&</sup>quot; NM = not measured.

 $<sup>^{\</sup>nu}$  NC = not calculated.

B ----- = not analyzed; MPE and MPF were not installed until July 1995.

bioventing at the site. Respiration and fuel biodegradation rates at MPF were somewhat elevated during the Option 1 testing event due to the remaining hydrocarbon contamination present at this location.

#### 2.4.4 Recommendations Following Option 1 Testing

Because of the relatively high TVH concentrations in soil gas at MPF, Parsons ES recommended continued expanded-scale bioventing system operation at the site for an additional 6 months to 1 year prior to initiating Option 2 confirmation sampling activities. Based on soil sampling results and soil gas and respiration results obtained following approximately 3 years of SVE and pilot- and expanded-scale bioventing system operation, the majority of the soils at the site are likely to have been sufficiently remediated to meet FDEP (1997) criteria outlined in FAC, Chapter 62-770. Continued operation of the expanded-scale bioventing system since November 1996 has further reduced petroleum hydrocarbon contamination in site soils, including soils near MPF. It is anticipated that soil analytical results will meet FDEP risk-based criteria for NFA, or groundwater monitoring only, and no further soil remediation will be necessary.

#### 2.5 NATURAL ATTENUATION UPDATE SAMPLING

In March 1998, personnel from NRMRL performed an additional round of groundwater sampling at Site ST-29 to evaluate the progression of dissolved fuel contaminant natural attenuation. At the time this SAP was prepared, monitoring results from this sampling event were not available. These groundwater sampling results will be provided in a separate report generated as part of the AFCEE natural attenuation project.

#### **SECTION 3**

#### SITE CLEANUP REQUIREMENTS

#### 3.1 SITE CHARACTERIZATION REQUIREMENTS

The objective of the confirmation sampling is to demonstrate that contaminant levels in soils contaminated by previous leaks of MOGAS from the fiberglass fuel line and UST located near the active fuel facilities west and southwest of Building 736 meet FDEP (1997) risk-based criteria for NFA, or groundwater monitoring only, and that no further soil remediation will be necessary. This SAP targets unsaturated and smear-zone soils in the immediate vicinity of, and downgradient from, the active BX Service Station pump islands and USTs.

#### 3.2 CLEANUP CRITERIA

This section describes Florida's closure approach for sites contaminated with petroleum products. The final draft *Petroleum Contamination Site Cleanup Criteria* rule (Chapter 62-770, FAC) (FDEP, 1997) presents guidance for determination of remedial requirements for closure of petroleum-contaminated sites, including several mechanisms for determining matrix-specific cleanup criteria. The regulations allow closure of petroleum release sites under several different scenarios, including:

- NFA Proposal Without Conditions,
- · NFA Proposal With Conditions, or
- Monitoring-Only Proposal for Natural Attenuation.

Closure of a site under the NFA-Without-Conditions alternative would allow unrestricted future use of the site (e.g., residential land use), and therefore the requirements and allowable contaminant levels under this alternative are the most restrictive. The NFA-With-Conditions alternative requires that appropriate institutional or engineering controls be implemented to limit receptor exposure to contaminated media; sites seeking closure under this alternative are subject to potentially less stringent cleanup levels. A natural attenuation monitoring program is a recognized means of remediating sites with petroleum hydrocarbon contamination in groundwater, with the goal of achieving the NFA target cleanup levels.

#### 3.2.1 No Further Action Without Conditions

Closure of a petroleum release site under an NFA Proposal without conditions requires that a site meet the following criteria:

- No mobile LNAPL (free-phase product) is present;
- No fire or explosion hazard is present due to release of petroleum or petroleum products;
- · No "excessively contaminated soil" is present; and
- Matrix-specific target cleanup levels (for soil and groundwater) are met.

Contaminant concentrations in all affected media at a site must be below all applicable target cleanup levels for the site to qualify for a NFA without conditions proposal. The *Petroleum Contamination Site Cleanup Criteria* rule (FDEP, 1997) incorporates matrix-specific target cleanup levels for petroleum constituents in the form of "look-up" tables or through reference to other applicable regulations (i.e., state groundwater or surface water regulations).

To demonstrate that contaminated soil is not present in the unsaturated zone, representative soil samples must show that concentrations of the applicable petroleum products' chemicals of concern are less FDEP (1997) direct human exposure and leachability target levels. If leachability target cleanup levels are exceeded, direct leachability testing can be performed to determine if leachate concentrations exceed the applicable groundwater target cleanup levels. In addition, the rule also allows for the development of alternative cleanup standards that can be used in place of those presented in the look-up tables. The alternative cleanup standards must be developed based on site-specific exposure scenarios and risk assessment.

#### 3.2.2 No Further Action With Conditions

Closure of a petroleum release site under an NFA Proposal with conditions requires that a site meet the first three criteria for NFA without conditions (Section 3.2.1); however, alternative target cleanup levels may be justified by the property owner by agreeing to the enactment of institutional controls (i.e., land use restrictions) and/or engineering controls. For soil, less restrictive direct-exposure target cleanup levels may be used, and leachability target cleanup levels may be exceeded if it can be demonstrated based on site characteristics and restrictions specified in the institutional control, that petroleum product chemicals of concern will not leach into groundwater at concentrations exceeding applicable groundwater target cleanup levels. For groundwater, alternative groundwater target cleanup levels may be justified depending on the current and projected use of groundwater in the vicinity and by enacting appropriate institutional controls.

#### 3.2.3 Natural Attenuation With Monitoring

The FDEP recognizes natural attenuation with monitoring as a viable site rehabilitation strategy for petroleum-contaminated groundwater. The following criteria must be met to demonstrate that this strategy is appropriate for a site:

No mobile LNAPL is present;

- Contaminated soil is not present or does not constitute a continuing source of contamination to groundwater;
- Groundwater contaminant concentrations above applicable target cleanup levels are not migrating beyond a temporary POC;
- Available data show an overall decrease in the mass of contamination; and
- Contaminant concentrations in groundwater do not exceed appropriate criteria (Table IX levels, Chapter 62-770, FAC); or technical evaluations (as specified in Chapter 62-770.690 (1)(f), FAC) indicate that natural attenuation is an appropriate remedial alternative.

Natural attenuation with monitoring requires the establishment of a temporary POC based on site-specific conditions relating to land and groundwater use, potentially exposed populations, hydrogeology, and type and concentrations of contaminants. If human health, public safety, and the environment are protected, the POC may be moved to the property boundary, or beyond the property boundary (with notice), if necessary to address current plume conditions.

#### 3.3 CLEANUP STANDARDS FOR THE BX SERVICE STATION, SITE ST-29

Based on previous soil sampling results and soil gas and respiration results obtained following approximately 3 years of SVE and pilot- and expanded-scale bioventing system operation, the majority of the soils at the site are likely to have been sufficiently remediated to meet FDEP (1997) criteria outlined in Chapter 62-770. Continued operation of the expanded-scale bioventing system since November 1996 has further reduced petroleum hydrocarbon contamination in site soils including soils near MPF. It is anticipated that the concentration of residual MOGAS chemicals of potential concern (COPCs) in unsaturated and smear zone soils in the vicinity of the active dispensing islands and USTs at the BX Service Station are less than the applicable FDEP target cleanup levels for NFA and no further soil remediation will be necessary.

#### 3.3.1 Soil Criteria

Confirmation soil sample results will be compared to target cleanup levels established by FDEP (1997) to demonstrate that petroleum hydrocarbon contaminants in site soils have been sufficiently remediated to meet the requirements for a NFA proposal. Specifically, direct-exposure and leachability target cleanup levels from Table IV (FDEP, 1997) will be used as the initial cleanup criteria. Confirmation soil sampling results will be compared to the residential and industrial direct-exposure levels and leachability target levels for groundwater resource protection and recovery (because shallow groundwater at the site is not likely to impact surface water in the vicinity of Patrick AFB).

Table 3.1 compares the maximum detected site soil concentrations of petroleum hydrocarbon contaminants to the proposed FDEP target cleanup levels. As is evident from this comparison, pre-bioventing concentrations of benzene, ethylbenzene, xylenes and TRPH in soil were not in compliance with FDEP (1997) direct-exposure or

#### TABLE 3.1 COMPARISON OF MAXIMUM SITE SOIL CONCENTRATIONS TO TARGET CLEANUP LEVELS

#### BX SERVICE STATION SITE ST-29 PATRICK AFB, FLORIDA

		Maximum	Location of	Sample	FDEP	Target Clea	nup Levels a/
		Concentration	Maximum	Collection	Direct E	xposure <sup>b/</sup>	Soil
Chemical Name	Units	Detected	Detection	Date	I	II	Leachability c/
TRPH d/	mg/kg e/	2,730	HVW-2	Mar-93	350	2,500	340
Benzene	mg/kg	6.99	CPT-02-A19	Mar-94	1.1	1.50	0.007
Ethylbenzene	mg/kg	260	HVW-2	Mar-93	240	240	0.4
Toluene	mg/kg	54	HVW-2	Mar-93	300	2,000	0.4
Xylenes	mg/kg	2,600	HVW-2	Mar-93	290	290	0.3
Acenaphthene	mg/kg	ND f/	NA g/	Oct-93	2,300	22,000	4
Acenaphthylene	mg/kg	ND	NA	Oct-93	1,100	11,000	22
Anthracene	mg/kg	ND	NA	Oct-93	19,000	290,000	2,000
Benzo(a)anthracene	mg/kg	ND	NA	Oct-93	1.4	5.1	2.9
Benzo(a)pyrene	mg/kg	ND	NA	Oct-93	0.1	0.5	7.8
Benzo(b)fluoranthene	mg/kg	ND	NA	Oct-93	1.4	5	9.8
Benzo (g,h,i)perylene	mg/kg	ND	NA	Oct-93	2,300	45,000	13,000
Benzo(k)fluoranthene	mg/kg	ND	NA	Oct-93	15	52	25
Chrysene	mg/kg	ND	NA	Oct-93	140	490	80
Dibenzo(a,h)anthracene	mg/kg	ND	NA	Oct-93	0.1	0.5	14
Fluoranthene	mg/kg	0.20J h/	SB-10	Oct-93	2,800	45,000	550
Fluorene	mg/kg	ND	NA	Oct-93	2,100	24,000	87
Indeno(1,2,3-cd)pyrene	mg/kg	ND	NA	Oct-93	1.5	5.2	28
Naphthalene	mg/kg	60	SB-15	Oct-93	1,000	8,600	
Phenanthrene	mg/kg	0.12J	SB-10	Oct-93	1,900	29,000	120
Pyrene	mg/kg	0.14J	SB-10	Oct-93	2,200	40,000	570
MTBE 1/	mg/kg	NR j/	NA	NA	350	6,100	0.2
Lead	mg/kg	15.3	SB-6	1990	500	1,000	100/TCLP k/

Notes: Shading indicates maximum site concentration above target cleanup level indicated.

<sup>&</sup>lt;sup>2</sup> FDEP, 1997.

<sup>&</sup>lt;sup>ы</sup> Direct Exposure I and II are for No Further Action Without or With Conditions.

 $<sup>^{\</sup>circ\prime}$  Based on Table V (Groundwater Cleanup Target Levels).

<sup>&</sup>lt;sup>th</sup> TRPH = total recoverable petroleum hydrocarbons. Maximum concentration based on analysis by USEPA Method 418.1; total petroleum hydrocarbons (TPH) concentrations from March 1994 not considered. Target cleanup levels based on State of Florida FL-PRO analysis.

e' mg/kg = milligrams per kilogram.

<sup>&</sup>quot; ND = not detected; laboratory reporting limit not available.

<sup>&</sup>lt;sup>g/</sup> NA = not applicable.

<sup>&</sup>lt;sup>by</sup> J = analyte detected below laboratory reporting limit; the value shown represents a laboratory estimated concentration.

W MTBE = methyl tert-butyl ether.

<sup>&</sup>lt;sup>j</sup> NR = no results available.

k/ If total lead concentration exceeds 100 mg/kg when analyzed by USEPA SW7421, then sample must pass Toxicity Characteristic Leaching Procedure (TCLP) criterion of 5.0 mg/L for total lead.

leachability target cleanup levels. However, it is anticipated that representative confirmation soil sample results will demonstrate that the concentration of these contaminants in soil following approximately 4.5 years of soil venting treatment now are below the prescribed levels.

#### 3.3.2 Alternative Cleanup Standards

For those contaminants exceeding target cleanup levels for NFA (With or Without Conditions), alternative cleanup standards for soil contaminated with petroleum products may be developed based on a site-specific risk assessment. Development of alternative standards for those chemicals that are below the more stringent levels provided in the Rule should not be necessary.

If performed, the risk assessment would include a site-specific exposure assessment based on:

- Chemical concentrations in all contaminated media;
- Soil properties;
- Potential exposure pathways and routes;
- Current or potential future exposed populations;
- Exposure factors (i.e., exposure duration and frequency); and
- Expected contaminant concentrations to which actual or potential receptors may be exposed.

In establishing alternative target cleanup levels, the following factors would be used, as appropriate:

- Calculations using a lifetime cancer risk level of 1.0 x 10<sup>-6</sup> for carcinogenic COPCs;
- A hazard index of 1 or less for noncarcinogenic COPCs:
- Best achievable detection limits;
- Naturally occurring background concentrations (for inorganics); and/or
- Nuisance, organoleptic, or aesthetic considerations.

These alternative site-specific target cleanup levels, if developed, would be used for closure under an NFA With Conditions proposal.

#### **SECTION 4**

#### SITE CONFIRMATION SOIL AND GROUNDWATER SAMPLING AND ANALYSIS PLAN

The following SAP describes the borehole locations, sampling depths, soil sampling procedures, and analytical methods proposed to collect sufficient data to verify remediation of petroleum hydrocarbon contaminants in vadose and smear zone soils at the BX Service Station to proposed cleanup levels (Table 3.1).

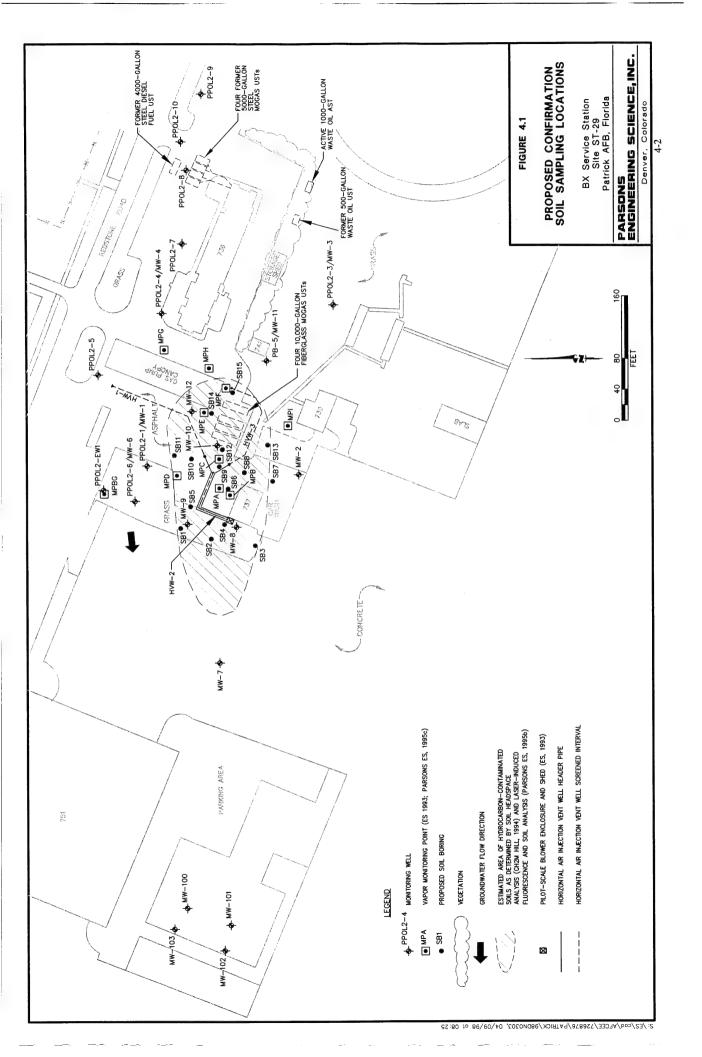
#### 4.1 SOIL SAMPLING

This section describes the scope of work required for collecting confirmation soil samples at the BX Service Station. An estimated 15 boreholes will be drilled and sampled in the vicinity of the active fueling facilities in the area previously shown to contain elevated hydrocarbon concentrations in saturated and unsaturated zone soils (Figure 2.6). A maximum of two additional boreholes may be drilled and sampled if field screening results indicate significant contamination extending beyond the proposed sampling area. Proposed borehole locations are shown on Figure 4.1.

One month prior to soil sampling, the expanded-scale bioventing system at the BX Service Station will be shut down to allow subsurface conditions to return to equilibrium. Soil sampling will be conducted by qualified Parsons ES scientists and technicians trained in the conduct of soil sampling, records documentation, and environmental sample chain-of-custody procedures. In order to provide complete documentation of the sampling event, detailed records will be maintained by the Parsons ES field hydrogeologist. In addition, sampling personnel will have thoroughly reviewed this SAP prior to sample collection and will have a copy available onsite for reference.

#### 4.1.1 Drilling, Sampling, and Equipment Decontamination

Soil boreholes will be advanced to the groundwater table surface (approximately 4 to 6 feet bgs) with a hand auger. Undisturbed soil samples, suitable for chemical analysis, will be obtained from each borehole by collecting the required volume of soil directly from the hand-auger bucket. Soil types will be classified according to the Unified Soil Classification System (USCS) and described in accordance with the standard Parsons ES soil description format. All soil samples will be visually examined, and sample headspace will be field screened for VOCs using a photoionization detector (PID) or a TVH analyzer (TVHA). The data obtained from the logging and screening will be recorded on borehole logs.



Based on field screening results, one sample with the greatest apparent petroleum hydrocarbon contamination from each boring will be selected and submitted for laboratory analysis using laboratory-prepared containers. Samples selected for laboratory analysis will be labeled with the site name and borehole number, sample depth, date of collection, project name, and other pertinent data. The sample containers will be sealed in plastic bags and immediately placed in an insulated cooler containing ice. The soil samples will be maintained in a chilled condition until delivered to the analytical laboratory. Chain-of-custody records will be prepared in the field and will accompany the samples to the analytical laboratory.

Augers and other downhole equipment will be cleaned before use and between boreholes to prevent cross-contamination. Between sampling events, the hand-auger bucket will be cleaned with Alconox® detergent, followed by successive potable and distilled water rinses. Decontamination water and auger cuttings will be managed as described in Section 4.4. Boreholes will be abandoned using bentonite following drilling and sampling. Boreholes drilled through asphalt will be repaired at the surface using asphalt cold-patch.

## 4.1.2 Soil Sample Analyses

Proposed soil sample analytical methods, estimated number of samples, and reporting limits are presented in Table 4.1. All samples will be analyzed by a State of Florida-certified and AFCEE-approved laboratory. Parsons ES proposes to analyze samples from the BX Service Station for TRPH by State of Florida Method FL-PRO; for BTEX using USEPA Method SW8020; and for PAHs using USEPA Method SW8310. Quality control (QC) samples also will be analyzed to assess laboratory methods. The laboratory will perform analyses on one matrix spike, one laboratory control sample, and one laboratory blank for each specific analytical method requested. Field QC samples will be collected and analyzed as described in Section 4.3.

### 4.2 CHAIN-OF-CUSTODY CONTROL

After the samples for laboratory analysis have been collected, chain-of-custody procedures will be followed to establish a written record of sample handling and movement between the sampling site and the laboratory. Samples collected for onsite field analyses will not require chain-of-custody records. Each shipping container will have a chain-of-custody form completed in triplicate by the sampling personnel. One copy of this form will be kept by the sampling contractor after sample delivery to the analytical laboratory, and the other two copies will be submitted to the laboratory with the samples. One of the laboratory copies will become a part of the permanent record for the sample and will be returned to Parsons ES with the sample analytical results. The chain-of-custody will contain the following information:

- Site name and address;
- Sample identification number;
- Sample collector's printed name and signature;

# TABLE 4.1 PROPOSED SOIL SAMPLE ANALYTICAL METHODS, REPORTING LIMITS, AND NUMBER OF SAMPLES

### BX SERVICE STATION

### SITE ST-29

### PATRICK AFB, FLORIDA

				Field or
	Number of	Reporting		Fixed-Base
Analytical Method	Samples <sup>a/</sup>	Limit <sup>b/</sup>	Units <sup>c/</sup>	Laboratory
State of Florida FL-PRO				
TRPH <sup>d'</sup>	15	TBD <sup>e/</sup>	mg/kg	Fixed-base
USEPA Method SW8020				
Benzene	15	1.0	μg/kg	Fixed-base
Toluene	15	2.0	μg/kg	Fixed-base
Ethylbenzene	15	2.0	μg/kg	Fixed-base
Xylenes	15	2.0	μg/kg	Fixed-base
MTBE <sup>ff</sup>	15	1.0	μg/kg	Fixed-base
USEPA Method SW8310				
Acenapthene	15	1,200	μg/kg	Fixed-base
Acenaphthylene	15	1,540	μg/kg	Fixed-base
Anthracene	15	440	μg/kg	Fixed-base
Benzo(a)anthracene	15	9	μg/kg	Fixed-base
Benzo(a)pyrene	15	15	μg/kg	Fixed-base
Benzo(a)fluoranthene	15	12	μg/kg	Fixed-base
Benzo(g,h,i)perylene	15	50	μg/kg	Fixed-base
Benzo(k)fluoranthene	15	11	μg/kg	Fixed-base
Chrysene	15	100	μg/kg	Fixed-base
Dibenzo(a,h)anthracene	15	20	μg/kg	Fixed-base
Fluoranthene	15	140	μg/kg	Fixed-base
Fluorene	15	140	μg/kg	Fixed-base
Indeno(1,2,3-cd)pyrene	15	30	μg/kg	Fixed-base
Naphthalene	15	1,200	μg/kg	Fixed-base
Phenanthrene	15	420	μg/kg	Fixed-base
Pyrene	15	180	μg/kg	Fixed-base

at Excludes QC samples. If optional boreholes are required, one additional soil sample per optional borehole also will be collected and analyzed.

<sup>&</sup>lt;sup>b/</sup> Project reporting limit as specified in subcontract for analytical services.

 $<sup>^{</sup>cl}$  mg/kg = milligrams per kilogram;  $\mu$ g/kg = micrograms per kilogram.

<sup>&</sup>lt;sup>d</sup> TRPH = total recoverable petroleum hydrocarbons.

e' TBD = to be determined.

<sup>&</sup>lt;sup>0</sup> MTBE = methyl tert-butyl ether.

- Date and time of collection;
- Place and address of collection;
- Type of sample (i.e. composite, grab, etc.);
- Sample matrix (soil or groundwater);
- Chemical preservatives added;
- Analytical laboratory to be utilized;
- · Analyses requested;
- Signatures of individuals involved in the chain of possession; and
- Inclusive dates of possession.

The chain-of-custody documentation will be placed inside the shipping container so that it will be immediately apparent to the laboratory personnel receiving the container, but will not be damaged or lost during transport. The shipping container will be sealed so that it will be obvious if the seal has been tampered with or broken.

### 4.3 QUALITY ASSURANCE/QUALITY CONTROL PROCEDURES

Samples must be collected, preserved, transported, and analyzed in such a manner that sampling results yield information that provides a reliable representation of the soil and groundwater quality at the site. To meet this requirement, the procedures described in Sections 4.1 will be followed during sample collection, handling, and analysis. In addition, laboratory QC samples will be analyzed as described in Sections 4.1.2.

Field quality assurance will include collection of field replicates, rinseate blanks, and trip blanks. Soil QC sampling will include two replicates (minimum frequency of 10 percent), one rinseate blank, and one trip blank for each cooler with samples designated for BTEX analysis.

### 4.4 MANAGEMENT OF INVESTIGATION-DERIVED WASTES

Soil cuttings generated during hand-auger soil sampling will be minimal. Residual cuttings not prepared for laboratory analysis will be returned to their respective soil borehole prior to plugging the borehole with bentonite. Equipment decontamination and rinseate water will be accumulated, transferred to a truck-mounted tank, and transported to the Trident STP (an industrial wastewater treatment plant) at Cape Canaveral Air Station (AS) for discharge and treatment. The site name, source location, volume, date of collection, and other pertinent information will be recorded in the Cape Canaveral AS investigation-derived waste inventory maintained by Parsons ES.

### SITE CONFIRMATION SAMPLING REPORT FORMAT

Following receipt and evaluation of the laboratory analytical results, a draft confirmation soil sampling report will be prepared. The report will summarize soil analytical results from the confirmation sampling event in order to demonstrate source removal and support recommendations for NFA (i.e., no further soil remediation), if appropriate. The confirmation sampling report and recommendations will be submitted to FDEP, Patrick AFB, and AFCEE.

As a minimum, the report will contain the following information for the BX Service Station:

- This confirmation SAP (as an appendix);
- Site plot plan showing sampling locations;
- Summary of field activities;
- Assessment of soil analytical results in comparison to applicable FDEP (1997) target cleanup levels (Tables 3.1);
- Laboratory analytical reports and chain-of-custody forms;
- Borehole logs;
- FDEP (1997) required information, conclusions, and recommendations for NFA (Chapter 62-770.680, FAC), natural attenuation and monitoring-only (Chapter 62-770.690, FAC), or additional active remediation (Chapter 62-770.700, FAC) or monitoring (Chapter 62-770. 750., FAC), as appropriate, based on available analytical results.

# PATRICK AFB SUPPORT REQUIREMENTS

The following Patrick AFB support is needed prior to the arrival of the Parsons ES team:

- Assistance in obtaining digging permits,
- · Arrangement of site access for Parsons ES, and
- Assistance in handling/disposal of decontamination/rinseate waters in accordance with Section 4.4.

# PROJECT SCHEDULE

The following schedule is contingent upon timely approval of this confirmation SAP and fulfillment of the Patrick AFB support requirements outlined in Section 6.

Event	Date
Submit Draft Confirmation SAP to AFCEE and Patrick AFB	29 April 1998
Receipt of AFCEE and Patrick AFB Comments	29 May 1998
Submit Final SAP to AFCEE, Patrick AFB, and FDEP*	June 1998
Confirmation Sampling	July 1998
Submit Draft Confirmation Sampling Report to AFCEE and Patrick AFB	October 1998
Receipt of AFCEE and Patrick AFB Comments	November 1998
Submit Draft Final Confirmation Sampling Report to AFCEE, Patrick AFB, and FDEP	December 1998

<sup>\*</sup> FDEP review and comment on the SAP occurred during the 17 April 1998 Petroleum Action Management Plan (PAMP) Meeting.

# POINTS OF CONTACT

Mr. Ed Worth 45 CES/CEV 1224 Jupiter Street Patrick AFB, FL 32925-3343 DSN 467-0965 COM (407) 853-0965 Fax: (407) 853-5435

Mr. Mike Deliz Florida Department of Environmental Protection 2600 Blair Stone Road, MS4505 Twin Tower Office Building Tallahassee, FL 32399-2400 (904) 921-9991

Major Ed Marchand AFCEE/ERT 3207 North Rd, Bldg. 532 Brooks AFB, TX 78235-5363 DSN 240-4364 COM (210) 536-4364 Fax: (210) 536-4330 Mr. Steve Archabal Site Manager Parsons Engineering Science, Inc. 2323 West 14th Street, Suite 616 Tempe, AZ 85281 (602) 921-0435 Fax: (602) 966-9119

Mr. John Ratz/Mr. Craig Snyder Project Manager/Deputy Project Manager Parsons Engineering Science, Inc. 1700 Broadway, Suite 900 Denver, CO 80290 (303) 831-8100 Fax: (303) 831-8208

Mr. Roger Bonner Parsons Engineering Science, Inc. 1485 S. Semoran Blvd., Suite 1450 Winter Park, FL 32792 (407) 671-5454 Fax: (407) 671-4199

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# APPENDIX A BX SERVICE STATION (SITE ST-29) GROUNDWATER DATA

# APPENDIX A-1 GROUNDWATER DATA FROM OB&G, 1994 AND 1995

USAF - 45th Space Wing Contamination Assessment Report ST-29 (PPOL-2) Revision: 0 November 9, 1994 Table 1-1

# TABLE 1-1 - ST-29 (PPOL-2) PREVIOUSLY DETECTED COMPOUNDS PHASE II, STAGE 2, ROUND 1 AND 2 GROUND WATER

			•									
	WELL PPOL2-1	POL2-1	WELL PPOL2-2	POL2-2	. WELL F	WELL PPOL2-3	WELL PPOL24	POL2-4	WELL PPOL2-5	POL2-5	WELL PPOL2-8	9012-8
Parameters Units	Stage 2 Round 1	Stage 2 Round 2										
Petroleum Hydrocarbons mg/L	0.8	-	_	-	1	ı	1	i	ı	1	ı	
Lead, lotal mg/L	0.0025	0.0101	-	0.0039	0.0070	0.0221	0.0017	0.0027	0.0033	0.0095	-	-
Lead, dissolved mg/L	١	-	. 1	-	ı	0.0029	ı	0.0014	1	1	-	-
Bromodichloromethane µg/L	1	-	t	ı	-	-	•	3.10	1	1		1
Chloroform	ı	_	-	1	Í	1	i	6.63	1	ı	ı	-
Chloromethane µg/L	1	-	669'0.	ı	QN	_	_	ı	ı	ı	1	1
Trans-1,2-dichloroethene µg/L.	1	0.31	١	1	-	1	1	ľ	t	0.65	-	1
Methylene Chloride	1	1	-	_	-	-	1	QN	,	ı	1	1
Trichloroethene µg/L	,	0.702	-	1	ı		_	lą.	-	ı	1	1
Chlorobenzene	0.387	1	1	-	_	1	1	1	ı	t	-	1
Toluene µg/L	0.85	٦	1	1	-	1	0.72	1	1	i	1	-
Xylenes, lotal µg/L	1.56	-	1	1	1	t	1.62	ı	١	ı	ı	1
Methyl-T-butylether µg/L	151	QN	0.910	ı	ı	1	13.3	1	,	QV	-	ı
Bis(2-ethythexyl) phthalate PGL	NRQ	-	NRQ	i	NRQ	ı	NRQ	ı	NRQ	1	NRQ	4.2

ND = KEY:

current concentration less than instrument detection limit Not Detected Not Requested

ESE, 1991. Source: PUP

PPOL2-10

PUP

PPOL2-

g

PPOL2-8

PPOL2-5

PPOL2-4

ST-29 (PPOL-2) Revision: 0 November 9, 1994 Table 2-6 (Page 1 of 5) 皇 9 g 읖 ž 9 9 9 2 2 ž 9 9 ž 9 皇 £ 9 œ 오 오 2 2 웆 2 皇 皇 울 읖 ž i 2 9 Ş 읖 g 9 9 9 9 ç 오 9 2 ž 2 2 ₹ 9 9 9 ž 웆 2 2 9 Round 1 Round 2 g 9 2 9 9 2 9 9 2 2 g 2 ¥ 웆 g 9 9 읒 운 9 물 2 무 읒 9 2 ž 2 Units Round 1. Round 2. Round 1. Round 2. Round 1. Round 2. Round 1. Round 1. Round 2. Round 1. Round 2. 2 2 2 9 9 9 ç 2 9 9 ç 9 웊 皇 皇 읒 2 ž 2 9 2 皇 ç 9 웊 웆 8 g 2 2 9 읖 웆 웆 웆 g 9 皇 S 9 8 S 2 ž 皇 웆 2 2 9 ç 읒 웊 g g 문 9 웆 Š 9 g ž 윷 읒 2 £ 2 皇 읖 9 읒 옃 9 2 2 9 2 웆 9 9 2 윷 2 9 9 9 9 身 ž 읒 皇 S 읖 2 웊 2 2 皇 2 9 웆 웆 皇 ŝ 9 읒 2 9 웊 웆 웆 9 2 웆 읖 2 ç 2 2 읒 9 ž 2 문 2 9 9 2 2 문 문 9 9 £ 문 웆 2 웆 2 웆 웆 2 웆 9 ž 皇 呈 읒 웆 皇 皇 2 읖 윷 9 2 9 웊 2 皇 용 웆 9 9 2 2 g g S g 9 2 웆 뎦 皇 S 읖 ĝ 2 읒 9 물 2 皇 9 S 웃 ž 呈 皇 읖 읖 윷 皇 9 9 2 2 9 9 身 2 g 읒 2 2 읖 ž 9 Ş 2 읖 2 9 皇 읖 2 呈 읖 2 9 9 9 £ 2 2 2 2 2 2 웆 2 皇 ž 2 - 9 2 2 2 9 웆 皇 9 웆 웆 VOLATILE ORGANICS (USEPA METHOD SW-846 8010/8020) 웆 S 2 오 £ 9 2 2 ₽ 2 웆 呈 웆 ç 2 g 9 2 呈 ž 웃 2 9 呈 皇 문 皇 9 2 2 g 웆 2 2 皇 9 웆 呈 2 2 2 ž 皇 2 읖 2 9 9 읟 2 2 2 9 9 呈 읒 身 ng/L ng/L 1/Bn ng/L ng/L 7/Bn 7/Bn ng/L ng/L 1/Bn ng/L ng/L ٦ N ng/Ľ 7/Bn ng/L ng/L ng/L ng/L 7/gn √Bn ng/F 7/Bn ug/L √g V ng/F ng/L ng/F 1,1,1,2-Tetrachloroethane 1,1,2,2-Tetrachloroethane trans-1,2-Dichloroethylene Dichlorodifluoromethane cls-1,3-Dichloropropane 2-Chloraethyl vinyl ether Dibromochloramethane trichlorofluoromethane Bromodichloromethane cis-1,2-Dichloroethene 1,2-Dichloropropana 1,2-Dichlorobenzena 1,4-Dichlorobenzene Carbon tetrachloride 1,3-Dichlorobenzene 1,1-Dichloroethane 1,1-Dichloroethene I,2-Dichloroethane Dichloromethane Dibromomethane 1-Chlorohexane Chloromethane 2-Chlorotoluene Bromobenzene Benzyl chloride Bromomethane Chlorobenzene Chloroethane PARAMETER Bromoform Chloroform

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PARAMETER. Tetrachloroethylene	Units	Units Round 1 Round 2	6 P C	•	6 60.00		6 PC					7	Round 2	, 	
strachioroethylene			Honina e	Round 1	מחוות ליי	Konna I konna z	House &	Hound 1 Hound 1	٠l	Honua z	u piunou	Honua z	1	Hound 1	
TO 10.1000 100 100 10 10 10 10 10 10 10 10 10	ng/L	QN	QN	8	QN	Q	S	Q.	QN	Q	Q	Q	Q	Q	Q
1,1,1.Trichloroethane	7/Bn	QN	9	2	2	9	Q	Q	2	9	2	9	Q	9	2
1,1,2-Trichloroethane	ng/L	Q	Q	2	QN	Q	9	Ω	9	Q	Q	Q	Q	Q	S
Trichloroethylene	ug/L	2	2	9	9	9	9	QV	9	2	2	9	9	9	2
1,2,3-Trichloropropane	ng/L	Q	QN	Q	Q	Q	2	Q	Q	Q	Q	QN	Q	Q	S
Vinyl Chloride	J/Bn	2	9	9	QN	9	9	Q	9	QV	Q	Q	9	9	2
trans-1,3-Dichloropropylene	ng/L	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	S
Ethylbanzena		g	2	2	g	2	9	8	4	43	웆	Q	Q	Q	2
Toluene	ng/L	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	QN	Q	Q	S
Xylenes, (Total)	T/Bn	9	Q	<b>(</b> ₹	QV /	9	9	Q	2	9	2	2	2	QN	2
Methyl tert-butyl ether	ng/L	Q	Q	15	QN	Q	Q	Q	Q	Q	QN	QN	Q	Q	Q
Total VOA	ng/L	0.0	0.0	) 55	0'0	1	0.0	41	7	68	0.0	0.0	0.0	0.0	0.0
Semi-Volatite Organics (USEPA Method SW846 8270) Acenaphthene ug/L ND NE	cs (USEPA Method SWB. ug/L ND	od SW84	6 6270) ND	Q	QN	QN	Q	QN	αN	QN	QN	Q	Q	g	g
ĺ		2	Ş	CZ	2	Ç	Š	5	Ç	Š	Ş	Ş	Ş	CM	Ş
Anthracene	J/Bn	2	2		2	2	2	2	2	2	2	2	9	2	9
Benzidine	ng/L	g	Q	g	Q	Q	Q	2	Q	g	g	Q	QN	Q	S
Benzo(a) anthracene	7/Bn	QN	Q	9	QN	ON	2		9	2	9	S	9	ž	<b>≨</b>
Benzo(b)fluoranthene	ng/L	Q	N	g	Q	QN	Q	Q	Q	Q	Q	Q	Q	Q	S
Benzoic Acid	1/Bn	N	Ą	Ϋ́	ž	NA A	۸N	¥	NA	¥	¥.	Ϋ́	Ą	Q	2
Benzo(k)fluoranthene	ug/L	Q	Q	QN	Q Q	S	S	Q	QN	Q	Q	QN	QN	QN	Q
Benzo(ghi)parylene	ng/L	9	2	QN	QN	QV	Q	2	Q	2	Q	Q	Q	Q	2
Benzo(a)pyrene	ng/L	ă	Q	Q.	ð	Q	Q	Q	2	Q.	2	Q	9	₽	S
Benzyl alcohol	ng/L	¥	¥	ď	ΝΑ	Ą	¥	¥	ž	Š	ž	ΥN	Ä	QN	2
Bis(2-chloroethoxy)methane	e ug/L	Q	å	Q	Q	Q	Q.	9	2	Q	Q	2	Q	2	S
Bis (2-chlaroethyl) ether	7/Bn	9	呈	g	QV	2	2	Q	9	2	2	2	QV	QV	2
Bis(2-chloroisopropyl)ether	r ug/L	S	S	Q	QN	Q	용	9	2	2	Q	Q	2	S	S
Bis (2-ethylhexyl) phthalate	ng/L	9	9	Q	Q	9	2		21	2	18	2	QN	Q	9
4-Bromophanyl phenyl ether	er ug/L	Q	Q Q	Q	·Θ	Q	2	9	Q	Q	Q	Q	QV	Q	S
												**************			

USAF - 45th Space Wing Contamination Assessment Report ST-29 (PPOL-2) Revision: 0 November 9, 1994 Table 2-6 (Page 3 of 5)

the same and the same of the s		•	Units Round 1 Round 2 R	ound 1	Round 2 R	Round 1 R	Round 2 R	Round 1 R	Round 1 R	Round 2 R	Round 1 R	Round 2 R	Round 2	Round 1	
PARAMETER	Units	Hound 1													
4-Chloroaniline	ng/L	¥ ¥	ž	¥	¥	Ą	Ą	¥	٧	ž	ž	¥	¥	2	2
4-Chloro-3-methylphenol	ng/L	2	9	Q	2	9	9	2	Q	Q	QN	2	Q	9	2
2-Chloronaphthalene	ng/L	QN	Q	g	Q	ð	Q	QN	Q	Q	Q	Q.	Q	2	S
2-Chlorophenol	ng/L	Q	2	9	Q	2	9	2	2	QN	2	2	QN	9	2
4-Chlorophenyl phenyl ether		9	Q	QN	Q	Q	QN	Q	Q	Q	S S	Q	9	Q	9
Chrysene		Q	Q	9	Q	9	QN	9	Q	QN	Q	QN	Q	QN	2
Di-n-butyl phthalate	ug/L	Q	Ş	Q	Q	QN	QN	2	Q	ND	Q	Q	Q	9	S
Dibenz(a,h)anthracene	ng/L	9	ž	2	NA	Q	Ϋ́	묏	9	ž	9	Ą	Ä	Q	2
1,2-Dichlorobenzene	ug/L	Q	Š	2	a	2	Q	S	Q	QN	Q	용	Q	Q.	2
1,3-Dichlorobenzene	J/Bn	QN	9	QV	QV	9	2	2	Q	9	9	2	2	QN	2
Dibenzofuran	ug/L	X A	٧	×	Ą	Ą	¥	ž	Ą	Ą	Y Y	Y Y	Ą	Q	S
3,3'-Dichlorobenzidine	ng/L	Q	2	2	Q	2	2	2	ON	2	2	Q	9	QN	2
2.4-Dichlorophenol	ng/L	2	2	ð	Q	9	2	Q	Q	Q	Q	Q	2	Q	S
Diethyl phthalate	ng/L	QN	QN	QN	QN	QN	QN	Q	S	QV	9	2	2	S	2
2,4-Dimethylphenol	ng/L	Q	운	QN	g	S	Q.	2	Q	Q	Q	QN	Q	Q	S
Dimethyl phthalate	7/Bn	QN	2	2	2	9	Q	2	ON.	ON	Q	2	Q	2	2
Di-n-octyl phthalate	ng/L	Q	2	N	Q	Q	Q	S	Q	Q	Q	g	QN	QN	Š
4,6-Dinitro-2-mehtylphenol	ng/L	2	2	2	9	2	2	2	Q	2	9	2	QN	QN	2
2,4-Dinitrophanol	ng/L	ð	2	Q	Q	Q	Q	2	Q	Q	Q	Q	Q	S	Q
2,4-Dinitratoluene	ng/L	9	9	QN	Q	9	2	Q	QV	9	9	2	Q	QN	2
2,6-Dinitrotoluene	ug/L	Q	2	Q	2	Q	Q	2	Q	Q	Q	2	2	Q	2
Fluoranthena	ng/L	2	2	2	Q	αN	9	Q	Q	2	9	2	QN	QV	2
Fluorena	ng/L	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	Q	QN	QN	S
Hexachlorobenzens	ng/L	Q	9	9	Q	Q	Q	Q	QN	Q	Q	9	Q	QN	Ş
Hexachlorobutadiene	ug/L	Q	S	Q	Q	Q	S	2	Q	Q	õ	2	Q	QN	Q
Hexachlorocyclopentadlene	ng/L	Q	2	2	9	2	9	2	ON	2	8	2	9	Q	2
Hexachloroethane	ug/L	QN	2	Q	QN	웆	Q	9	Q	2	Q	Q	Q	S	Š
Indeno(1,2,3-cd)pyrene	ng/L	Q	9	QN	QN	2	2	2	Q	9	2	2	2	Q	Ş
Isophorone	l/Bn	Q	S	2	QN	2	용	S	QN	R	Q.	Q	Q	Q	S
2-Methylnaphthelene	l/Bn	ž	ž	¥.	'Y	Ą	NA A	ž	NA	NA	NA	Ν	Š	QV	2
2-Methylphenol	l/Bn	¥.	ž	N N	¥	۸	٩	Ϋ́	¥ V	٧	Ϋ́	Ϋ́	ΑN	QN	S
4-Methylphenol	yon	Ϋ́	Y	ΨV	Ø.V	V N		3	3	914	VV				

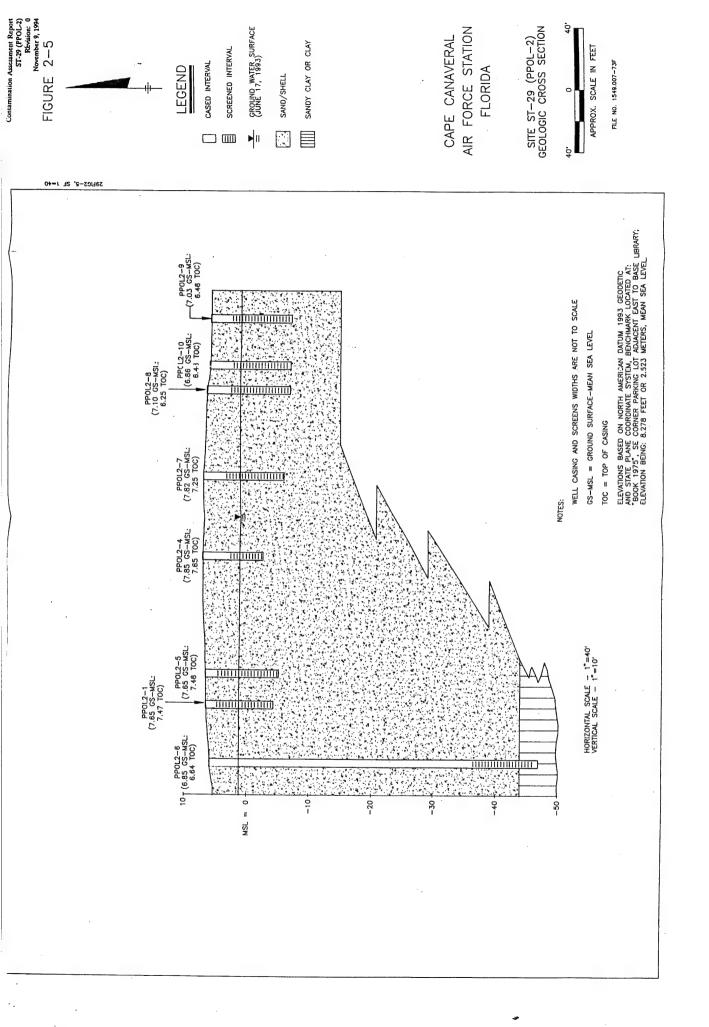
	<u> </u>	Pro12.4		י פאר פאר											
PARAMETER	Units A	lound 1	Units Round 1 Round 2	Round 1	Round 1 Round 2	Round 1	Round 2	Round 1	Round 1	Raund 2	Round 1	Round 2	Round 2	Round 1	
Naphthalene	l/Bn	S	S	Q	QN	QN	110	QN	87	Q.	Q	QN	Q	Q.	S
	VBn	ž	Ψ.	¥	ΝΑ	¥	¥	AA	ž	¥	ž	ž	Ą	Q	₹
Nitrobenzene	ug/L	문	Q	2	Q	Q	2	Q	Q	Q	Q.	Q	Q	2	2
3-Nitroaniline	/gn	ž	ž	₹	¥	Ϋ́	¥	₹	¥	¥	AN	¥	WA	Q	2
4-Nitroaniline	l/Bn	¥	¥	¥ X	¥	×	¥	¥	Ä	N A	¥	Y Y	N.	QN	9
2-Nitrophenal	ng/L	9	Q	2	2	9	2	9	Q	2	QN	g	2	Q	2
	ug/L	2	Q	Q	Q	Q	Q	Q	Q	Q.	QN	Q	Q	Q.	2
N-Nitrosodimethylamine	7/Bn	QV	2	2	Q	Q	2	2	9	9	Q	9	Q	Q	2
N-Nitrosodi-n-propylamine	ug/L	2	2	8	2	Q	Š	9	Q	QN	Q	Q	Q	Q	Q
N-Nitrosodiphenylamine	ng/L	QV	Q	Q	QN	2	2	2	9	9	9	2	9	9	2
Pentachlorophenol	ng/L	Q.	Q	Q	Q	Q	Q	Q	Q	Q	QN .	Q	Q	Q	2
Phenanthrene	1/Bn	2	2	2	2	2	2	9	2	QN	Q	9	QN	Q	9
Phenol	ng/L	2	Q	S	Q	Q	Q	Q	Q	Q	Q	2	ON	Q	S
Pyrene	ng/t	Q	QV	2	2	QN	2	2	2	9	2	9	Q	QN	2
1,2,4-Trichlorobenzene	ng/L	2	ð	g	2	QN	Q	Q	QN	Q	2	2	Q	Q	2
2,4,5-Trichlorophenol	l/Bri	Ϋ́	Ν	N	¥	NA	Ν	NA	NA	¥	ž	ž	٧	QN	Q
2,4,6-Trichlorophenol	ug/L	2	QN	Q	QN	Q	Q	Š	Q	Q	Q	Q	Q	٠Q	Q
1-Methylnaphthalene	l/Bn	ž	¥	ž	NA	¥	٧	ž	¥	≨	ž	ž	٧N	QN	2
Total PAHs (-Naphthalenes)	ng/L	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Table Manhallana	•	•	•	ď	•	ć			•						

TABLE 2-6 - ST-29 (PPOL-2) GHOUN" (TER SAMPLING ANALYSIS RESULTS

TABLE 2-6 - ST-29 (PPOL-2) GROUN" TER SAMPLING ANALTSISTIL SULTO

		PPOL24		PPO12-5	-	PPOL2.7	_	PPOLZ-8 DUP	ana		PPOL2-8		ana	DUP PPOL2:10 DUP	P.G.
PARAMETER	Units	Round 1	Round 2	Units Round 1 Round 2 Round 1 Round 2 Round 2 Round 1 Round 1 Round 2 Round 2 Round 2	Round 2	Round 1	Round 2	Round 1	Pound 1	Round 2	Round 1	Round 2	Round 2	Round 1	
LEAD (MCAWW 239.2)									2000						000000000000000000000000000000000000000
Total	J/Bn	10	ю	QN	4.	47	7	9	9	27	36.4	17	23	320	62
Dissolved	ng/L	8	ð	S	Q.	Q.	9	Q.	Q	8	2	Q	Q.	Q	2
(1818) METHOD EATED TO ME DATE OF THE WASHINGTON A 11 SEPA METHOD EA18		CBCXH	SNORG &	" SEPA M	FTHOO F4	. (181)									
TRPH	EINOLEU mg/L	mg/L ND	QN ND	ON ND	QN	αN	ON	QN	ON	QN	QN	QN	gN	Q	QN
Notes:	. ६					: .	·	:		:					
NA = Not Analyzed															
ND = Not detected															
DUP = QC/QA Duplicate Sample	Sample														
J = Estimated Value. Detected, but below quantitation limit.	tected, but	t below qu	antitation	limit											
Round 1 Ground Water Samples Collected June 17 through July 16, 1993 (PPOL2-5 resamples for SVOCs July 16, 1993 due to laboratory error.)	amples Co	elected Ju	ine 17 thro	ugh July 16	6, 1993 (P	POL2-5 re	samples f	or SVOCs	July 16, 1	1993 due	to laborato	ory error.)			
Round 2 Ground water samples collected August 1	amples col	lected Aug	gust 18 an	d 19, 1993	(PPOL2-4	resample	d for VOC	s, SVOCs,	TRPH, To	otal Disso	Ived Pb Oc	stober 8, 1	1993 due t	8 and 19, 1893 (PPOL2-4 resampled for VOCs, SVOCs, TRPH, Total Dissolved Pb October 8, 1893 due to laboratory error.)	/ error.)
Monitoring Well PPOL2-10 sampled for VOCs and SVOCs, January 1994; sampled for TRPH, Total Dissolved Pb. February 1994.	0 sampled	for VOCs	and SVOC	3s, January	r 1994; sar	npled for	TRPH, Tot	tal Dissolv	ed Pb. Fe	bruary 19	94.	٠			

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# APPENDIX A-2 GROUNDWATER DATA FROM PARSONS ES, 1995B

TABLE 4.3

# FUEL HYDROCARBON COMPOUNDS DETECTED IN GROUNDWATER, MARCH 1994 SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

	201 2025 202D 2038 2038 2038	Sample	Benzene	Toluene	Ethylbenzene	" Vulena	m. Yulene	, V. don	Total	Total	1 1 6 TM		E
	001 002S 002D 003S 003M	2 2 2	()(B/L)	(µg/L)	(HB/L)	p-Ayiene (µR/L)	(ILB/L)	O-Aylene (Hg/L)	Nytenes (µg/L)		(µg/L)	(µg/L) (µg/L)	1,2,3-TMB (µg/L)
	001 002S 002D 003S 003M												
	002D 002D 003M 003D	3-23-94	BLQ"	2.42	BLQ	BLQ	BLQ	BLQ	пго	2.42	91.0	BLQ	BLQ
	002D 003S 003M 003D	3-23-94	375	18.9	165	166	353	119	638	1196.9	71.2	NAW	86.3
	003K	3-23-94	1.5	1.7	1:1	1.4	2.6	1.4	5.4	9.7	δΊЯ	NA	1.3
7 3 3	003D	3-23-94	724	737	823	1220	2.110	1390	5020	7304	347	NA	403
	303D	3-23-94	207	15.6	40.5	42.2	24	7.5	73.7	336.8	2.8	NA	91
	2000	3-23-94	1.8	1.1	BLQ	BLQ	1.4	BLQ	1.4	4.3	91.0	٧٧	MQ
	0045	3-23-94	BLQ	9	DI.Q	ЫQ	) OIII	DTG	ेगा	9	ВГО	BLQ	131.Q
	0041)	3-25-94	BLQ	3.7	BI.Q	BLQ	ыго	BLQ	ЫQ	3.7	BLQ	DLQ	IILQ
	0058	3-25:94	BLQ	1.2	DI.Q	131.Q	131.Q	ભાવ	131.Q	1.2	ыго	NA	BLQ
	S900	3-25-94	BLQ	11.1	BLQ	. 1.3	2.3	рго	3.6	14.7	ВГО	٧×	BLQ
	SZ 00	3-25-94	BI.Q	3.9	ВГQ	BLQ	_	BLQ	_	4.9	BLQ	NA	BLQ
	S800	3-25-94	BLQ	2.8	BLQ	BLQ	BLQ	BLQ	ВГО	2.8	BLQ	BLQ	BLQ
	S600	3-25-94	2	4	BLQ	BLQ	ðп	ВГО	BLQ	9	BLQ	910	BLQ
	0000	3-25-94	427	14.1	2.9	11.7	12.1	6.6	33.5	477.5	ВГО	NA	9.4
	S010	3-25-94	nro	3.1	BLQ	BLQ	δηн	BLQ	дго	3.1	ВГО	BLQ	BLQ
$\sqcap \uparrow$	0118	3-25-94	BLQ	-	BLQ	BLQ	дго	BLQ	BLQ	_	BLQ	BLQ	BLQ
T	0125	3-25-94	BLQ	=	BLQ	ыд	1.2	BLQ	1.2	2.3	ЭГО	NA	BLQ
	012D	3-25-94	93.5	5.9	DI.Q	8.4	7	3.7	19.1	118.5	рц	NA	4.1
٦	0138	3-25-94	BLQ	8.4	) (10	• 10.4	5.4	3.1	18.9	\$6.4	ВГQ	ΝA	6.7
$\neg$	S910	3-26-94	-	1.9	ыго	DI.Q	BLQ	BLQ	оля	2.9	рпо	DI.Q	BLQ
	016DD	3-31-94	BI.Q	1.9	ысд	BLQ	ЫQ	131.0	BLQ	1.9	ыд	BLQ	BLQ
- 1	018S	3-24-94	BI.Q	1.7	2.3	3.8	9.9	4	14.4	18.4	1.4	NA	1.6
CPT-18D CPT-86-018D	018D	3-24-94	8.3	2.1	BLQ	BLQ	BLQ	ыго	BLQ	10.4	ВГО	вго	ЫQ
	018DD.	3-31-94	BI.Q	3.5	BLQ	BLQ	BI.Q	BLQ	пго	3.5	ВІ.Q	BLQ	ВГО
1	0198	3-24-94	BLQ	BI.Q		1.5	2.7	вго	4.2	5.3	ыд	NA	ВГО
	019D	3-24-94	1.5	BLQ	BLQ	BLQ	вго	BLQ	пл	1.5	ВГО	DLQ	вго
$\neg$	020S	3-24-94	BLQ	2.1	BLQ	1.5	2.7	1.8	9	8.1	δΊШ	NA	-
	020D	3-24-94	BLQ	BIQ	BLQ	вго	• бав	BLQ	òna	BLQ	рго	BLQ	BLQ
	021S	3-25-94	NA.	5.1	BLQ	BLQ	δΉθ	BLQ	δπя	5.1	DГО	٧V	BLQ
	021D	3-25-94	NA	BLQ	BLQ	ВГО	δηα	BLQ	ВГQ	BLQ	BLQ	BLQ	BLQ
	022S	3-26-94	NA	1.9	BLQ	BLQ	вго	BLQ	BLQ	1.9	<b>д</b>	BLQ	BLQ
	022D	3-26-94	1.5	BLQ	вго	BLQ	อาย	BLQ	BLQ	1.5	BLQ	BLQ	BLQ
	0238	3-25-94	BLQ	δΊЯ	91.0	ВГО	1.7	BLQ	1.7	1.7	BLQ	NA	131.Q
	023D	3-25-94	BLQ	BLQ	BLQ	BLQ	BLQ,	BLQ	BLQ	BLQ	δπа	BLQ	BLQ
CPT-24S   CPT-86-024S	024S	3-26-94	BLQ	BLQ	BLQ	DI.Q	B1.Q	BLQ	BLQ	BLQ	BLQ	BLQ	BLQ

# TABLE 4.3 (CONCLUDED)

# FUEL HYDROCARBON COMPOUNDS DETECTED IN GROUNDWATER, MARCH 1994 SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

Sample         Sample         Benzene         Toluene         Ethylbenzene         p-Xylene         m-Xylene         o-Yylene           Number         Date         (µg/L)										Total	Total			
ion         Number         Date         (µg/L)	Sample	Sample	Sample	Benzene	Toluene	Ethylbenzene	p-Xylene	m-Xylene	o-Xylene	Xylenes	BTEX	1,3,5-TMB	1,3,5-TMB 1,2,4-TMB 1,2,3-TMB	1,2,3-TMB
24D         CPT-86-024D         3-26-94         BLQ         I.4         6.9           25D         CPT-86-025B         3-26-94         BLQ         I.1         BLQ         BLQ         I.4         6.9           26S         CPT-86-026B         3-26-94         BLQ         I.1         BLQ	Location	Number	Date	(µg/L)	(µg/L)	(µg/L)	(µg/L)	(µg/L)	(µg/I.)	(µg/L)	(µg/L)	(μg/L)	(µg/I.)	(µg/L)
24D         CPT-86-024D         3-26-94         BLQ														
25S         CPT-86-025S         3-26-94         BLQ         2.1         2.3         4.1         6.9           25D         CPT-86-025D         3-26-94         BLQ         1         BLQ         BLQ         1.4           26S         CPT-86-026S         3-26-94         BLQ         1.1         BLQ         BLQ         BLQ           26D         CPT-86-026D         3-26-94         BLQ         1.1         BLQ         BLQ         BLQ           26D         CPT-86-026D         3-26-94         BLQ         1.1         BLQ         BLQ         BLQ           14D         CPT-86-014D         3-25-94         960         16.6         11.5         39.2         36.8           14D         CPT-86-010         3-23-94         4.1         BLQ         BLQ         1.1         1           101         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ           102         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ           103         CPT-86-102         3-23-94         BLQ         BLQ         BLQ         BLQ           103         CPT-86-102         3-23-94         BLQ         BLQ </td <td>CPT-24D</td> <td>CPT-86-024D</td> <td>3-26-94</td> <td>BLQ</td> <td>BLQ</td> <td>BLQ</td> <td>BLQ</td> <td>BLQ</td> <td>BLQ</td> <td>DTE</td> <td>BLQ</td> <td>BLQ</td> <td>BLQ</td> <td>BLQ</td>	CPT-24D	CPT-86-024D	3-26-94	BLQ	BLQ	BLQ	BLQ	BLQ	BLQ	DTE	BLQ	BLQ	BLQ	BLQ
25D         CPT-86-025D         3-26-94         BLQ         I         BLQ         II.4         II.4           26S         CPT-86-026S         3-26-94         BLQ         I.1         BLQ         BLQ         BLQ           26D         CPT-86-026D         3-26-94         BLQ         I.1         BLQ         BLQ         BLQ           14D         CPT-86-014D         3-25-94         BLQ         11.5         39.2         36.8           100         CPT-86-100         3-23-94         BLQ         BLQ         BLQ         BLQ           101         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ           102         CPT-86-102         3-23-94         BLQ         BLQ         BLQ         BLQ           103         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ           103         CPT-86-102         3-23-94         BLQ         BLQ         BLQ         BLQ           103         CPT-86-103         3-23-94         BLQ         BLQ         BLQ         BLQ           103         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ           103 <td>CPT-25S</td> <td>CPT-86-025S</td> <td>3-26-94</td> <td>BLQ</td> <td>2.1</td> <td>2.3</td> <td>4.1</td> <td>6.9</td> <td>3.5</td> <td>14.5</td> <td>18.9</td> <td>1.7</td> <td>N.A</td> <td>1.5</td>	CPT-25S	CPT-86-025S	3-26-94	BLQ	2.1	2.3	4.1	6.9	3.5	14.5	18.9	1.7	N.A	1.5
26S         CPT-86-026S         3-26-94         BLQ         1.1         BLQ         BLQ         BLQ           26D         CPT-86-026D         3-26-94         BLQ         1         BLQ         BLQ         BLQ           14D         CPT-86-04D         3-25-94         960         16.6         11.5         39.2         36.8           100         CPT-86-00         3-23-94         4.1         BLQ         BLQ         BLQ         BLQ           101         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ           102         CPT-86-102         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ           103         CPT-86-103         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ           103         CPT-86-103         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ           103         CPT-86-103         3-23-94         BLQ         BLQ </td <td>CPT-25D</td> <td>CPT-86-025D</td> <td>3-26-94</td> <td>BLQ</td> <td>1</td> <td>BLQ</td> <td>BLQ</td> <td>1.4</td> <td>BLQ</td> <td>1.4</td> <td>2.4</td> <td>BLQ</td> <td>NA</td> <td>BLQ</td>	CPT-25D	CPT-86-025D	3-26-94	BLQ	1	BLQ	BLQ	1.4	BLQ	1.4	2.4	BLQ	NA	BLQ
26D         CPT-86-026D         3-26-94         BLQ         I         BLQ         BLQ         BLQ           14D         CPT-86-014D         3-25-94         960         16.6         11.5         39.2         36.8           100         CPT-86-010         3-23-94         4.1         BLQ         BLQ         1.1         1           101         CPT-86-015         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ           102         CPT-86-101         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ           103         CPT-86-103         3-23-94         BLQ         BLQ         BLQ         BLQ         BLQ         BLQ           103         CPT-86-103         3-23-94         BLQ	CPT-26S	CPT-86-026S	3-26-94	ыго	1.1	ВГО	BLQ	DГО	BLQ	BLQ	1.1	BLQ	BLQ	BLQ
14D   CPT-86-0141)   3-25-94   960   16.6   11.5   39.2   36.8   11.0   CPT-86-100   3-23-94   4.1   BLQ   BLQ   I.1   1   1   1   1   1   1   1   1   1	CPT-26D	CPT-86-026D	3-26-94	BI.Q	1	BLQ	BLQ	BLQ	BLQ	BLQ	1	BLQ	BLQ	BLQ
100   CPT-86-100   3-23-94   4.1   BLQ   BLQ   1.1   1   1	CPT-14D	CPT-86-0141)	3-25-94	960	16.6	11.5	39.2	36.8	44.2	120.2	1108.3	15.3	NA	23
CPT-86-P135 3-26-94   BLQ   BLQ	NIW-100	CPT-86-100	3-23-94	4.1	BLQ	BLQ	1.1		BLQ	2.1	6.2	BLQ	BLQ	BLQ
CPT-86-101 3-23-94 BLQ BLQ BLQ BLQ BLQ CPT-86-102 3-23-94 BLQ BLQ BLQ BLQ BLQ CPT-86-103 3-23-94 BLQ BLQ BLQ BLQ BLQ CPT-86-P01.2-6 3-26-94 BLQ	PBS	CPT-86-PB5	3-26-94	BLQ	BLQ	91.0	BLQ	BLQ	DI O	BI.Q	BLQ	BLQ	BLQ	BLQ
CPT-86-102 3-23-94 BLQ BLQ BLQ BLQ BLQ BLQ CPT-86-103 3-23-94 BLQ BLQ BLQ BLQ BLQ CPT-86-PPOL2-6 3-26-94 BLQ	NIW-101	Cl <sup>v</sup> F-86-101	3-23-94	131.Q	BLQ	91.0	BLQ	DIEQ.	DI.Q	BLQ	BLQ	ыд	DI.Q	BLQ
CPT-86-103 3-23-94 BLQ BLQ BLQ BLQ BLQ BLQ CPT-86-PPOL2-6 3-26-94 BLQ	MW-102	CPT-86-102	3-23-94	BLQ	BLQ	BI.Q	BLQ	BLQ	BLQ	BLQ	BLQ	BLQ	BLQ	BLQ
CPT: 86-PPOL2-6 3-26-94 BLQ BLQ BLQ BLQ BLQ BLQ	MIW-103	CPT-86-103	3-23-94	BLQ	ыд	ЫQ	BLQ	ЫQ	BLQ	BLQ	) SIS	91.0	131.Q	BLQ
Fit Old Old Old Old Post Fit long stand	PPOI.2-6	CPT-86-PPOL2-6	. 3-26-94	BLQ	BLQ	BLQ	BI.Q	DI O	DTG	BLQ	BI.Q	OH	DLQ	DI Q
CT 1-80-1 CLZ-1 3-28-54   DLQ   DLQ   DLQ   LA	PPOL2-1	CPT-86-PPOL2-1	3-26-94	BLQ	- BLQ	BLQ	BLQ	1.4	<u></u> для	1.4	1.4	дп	ÒΊΘ	BLQ

" BLQ = Below quantitation limit, or < 1  $\mu$ g/L. " NA=Not available.

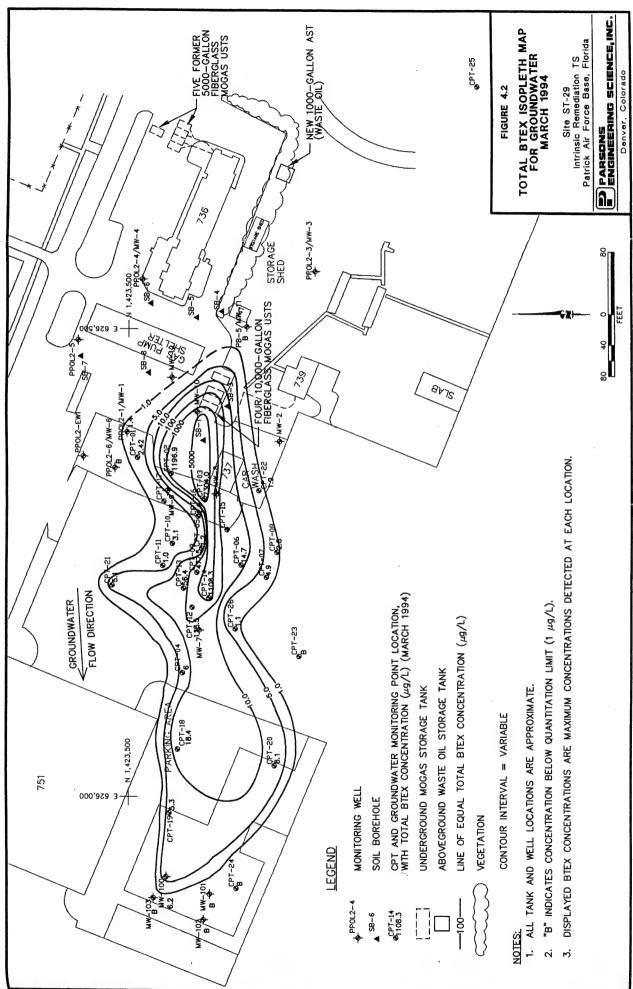
# FUEL HYDROCARBON COMPOUNDS DETECTED IN GROUNDWATER, MARCH AND MAY 1995 SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

	r								$\neg$		$\neg$													_	_	$\neg$					_		1	
TOC (mg/L)		14.9	16.3	13.8	NA™	35.8	181.9	21.8	3	10.9	6.1	9.5	3.4	7.8	7.5	7.5	4.6	5.8	4.2	4	3.3	Ϋ́	4.9	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
Total TMB (µg/L)	, ,	336.53	340.45	6.24	6.32	57	4530.83	10.09	7.15	5.23	1.27	1.04	2.37	3.09	3.11	3.12	ND	ND	ND	ND	ND	UN	ND	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
1,2,3 TMB (µg/L)	7 7 7	81.68	16.68	2.87	3.26	18.58	850.57	7.26	3.6	2.75	1.27	ND	ND	1.98	1.84	1.42	ND	UN	ND	ND	GN	CIN	ON	٧٧	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
1,3, 5 TMB 1,2,4 TMB 1,2,3 TMB (ug/L) (ug/L)		203.4	202.48	1.97	1.64	30.67	2903.48	1.45	2.46	1.33	CIN	ND	1.31	ND	ND	1.7	ND	ND	ND	ND	ND	ND	ND	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
1,3, 5 TMB (µg/L)	, A	43.95	48	1.4	1.42	7.75	81.911	1.38	1.09	1.15	CIN	1.04	1.06	1.11	1.27	ON	ON	QN	DN	QN	QN	ND	ON	NA	٧٧	NA	NA	NA	NA	NA	NA	NA	ΝΑ	NA
Total BTEX (ug/L)	7 0 1	566.4	583.21	68.28	62.64	292.99	14095.76	64.29	7.38	3.53	1.08	96'0	1.67	4.54	4.55	66'0	1.03	BLQ	ND	ON	ND	ND	ND	GN	ON	ND	ND	ND	ND	2945	83	160	CIN	3220
Total Xylene (tig/L.)	7 8 7	260.27	284.99	26.84	20.95	122.75	8820.63	14.18	5	3.53	1.08	96.0	PLQ	3.54	3.58	0.99	1.03	BLQ	ND	ND	ND	ND	UN .	, ND	ND	ND	ND	ND	ND	2450	ND	220	ND	2310
o-Xylene (ng/L)	7 8 7	8.58	9:36	1.39	2.05	32.14	2498.73	5.36	1.45	0.09	ЫQ	ND	CIN	1.35	1.3	ND	BLQ	GN	QN	UN	CIN.	QN	QN	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA
m-Xylene	7-8-1	156.04	168.18	2.96	1.12	32.43	3466.78	4.01	1.73	1.33	1.08	96'0	bгó	1.13	1.22	66.0	1.03	BLQ	ND	CIN	CIN	ND	CIN	NA	NA	NA	NA	NA	NA	NA	NA	NA	VN .	· NA
p-Xylene (119/L)	75 8 57	95.65	107.45	22.49	17.78	58.18	2855.12	4.81	1.82	1.21	ВГО	BLQ	BLQ	1.06	1.06	BLQ	BLQ	BLQ	ND	UN	GN	GN	GN	NA	VN	NA	NA	NA	NA	NA	NA	NA	NA	۲×
Ethylbenzene (ug/L)	/- 0.JV	132.59	135.4	10.49	10.39	21.8	2252.51	96.0	1.29	BLQ	DIG	BLQ	GN	BLQ	ВГО	BLQ	BI.Q	NI)	UD	ON	CIN	ND	ND	ND	ND	ND	QN	GN	OIN	270	46	360	ON	700
Tolucne (ug/L)	7.0.7	6.25	6.37	BLQ"	0.99	14.03	1526.23	4.35	BLQ	BLQ	ND°	GN	1.67	-	76.0	QN	DI O	CIN	ΩN	QN	QN	GN	GN	GN	CIN	GN	QN	QN	ND	8.5	ON	QN	CIN	QN
Benzene (ue/L)	/2.62/	167.29	156.45	30.95	30.31	134.41	1496.39	44.78	1.09	BLQ	BLQ	BLQ	BLQ	91.0	BLQ	BLQ	BLQ	ON	ND	ND	GN	ND	ND	CIN	ND	CIN	ND	ND	ND	140	37	180	UN	210
Sample		3/23/95	3/23/95	3/23/95	3/23/95	3/23/95	3/23/95	3/25/95	3/25/95	3/25/95	3/25/95	3/26/95	3/26/95	3/25/95	3/23/95	3/23/95	3/31/95	3/24/95	3/31/95	3/24/95	3/23/95	3/25/95	3/25/95	\$/95	\$6/\$	5/95	5/95	5/95	\$/95	\$/95	5/62	5/95	5/95	5/95
Sample		2S	2S DUP	3D	3D DUP	3M	38	9D	· \$6	12D	12S	26D	26S	86-4I)	86-4S	86-4S DUP	86-16DD	86-MW18D	36-MW18DD	86-MW18S	86-MW100	86-MW21D	86-MW21S	•	•	6	•			•	•		•	•
Sample		CPT-02S 2	Г	CPT-03D 3		CPT-03M 3	CPT-03S 3	CPT-09D 9	CPT-09S 9	CPT-12D 1		CPT-26D 2	CPT-26S 2	CPT-04D 8	CPT-04S 8	CPT-04S 8	CPT-16DD 8	CPT-18D 8	3PT-18DD 8	CPT-18S 8	MW-100	CPT-21D 8		MM-1ª	MW-2 <sup>4</sup>	MW-3 <sup>d</sup>	MW-4	<i>γ</i> 9-ΜΜ	™-7°	/4W-8 <sup>4</sup> /	<sub>\$</sub> 6-₩₩	™W-10	MW-11 <sup>4</sup>	√W-12 <sup>4</sup>

at BLQ = Below limit of quantitation, 1.0 µg/L.

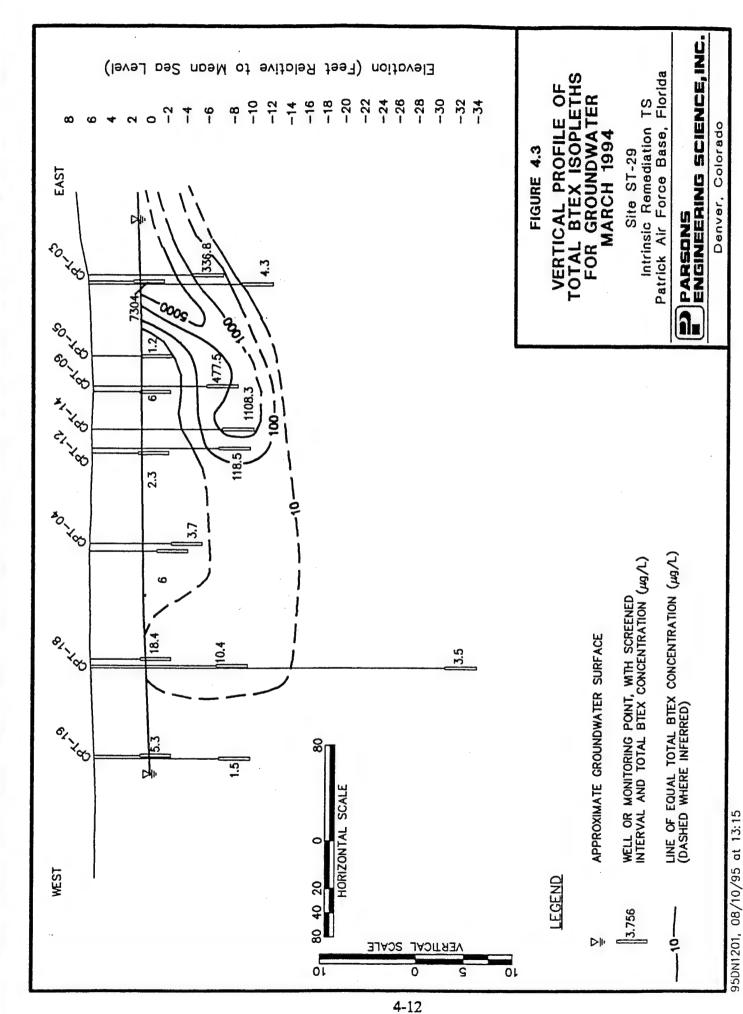
b/ NA = Data not available or sample not analyzed for this parameter.

c/ ND = Compound not detected at the method detection limit.
1/ Sample collected and analyzed by CII2M Hill on May 10 and 11, 1995.



950N1195, 08/10/95 at 11:00

4-11



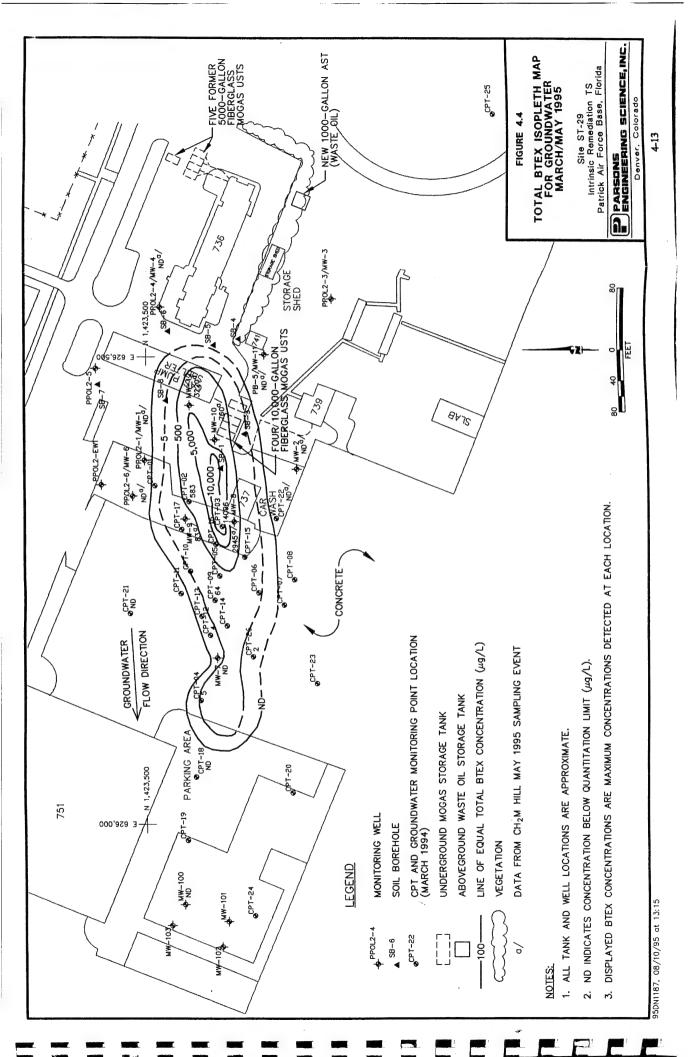


FIGURE 4.5A

# PLOT OF TOTAL BTEX VERSUS TIME FOR SAMPLING LOCATIONS WITH DECREASING BTEX CONCENTRATIONS SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

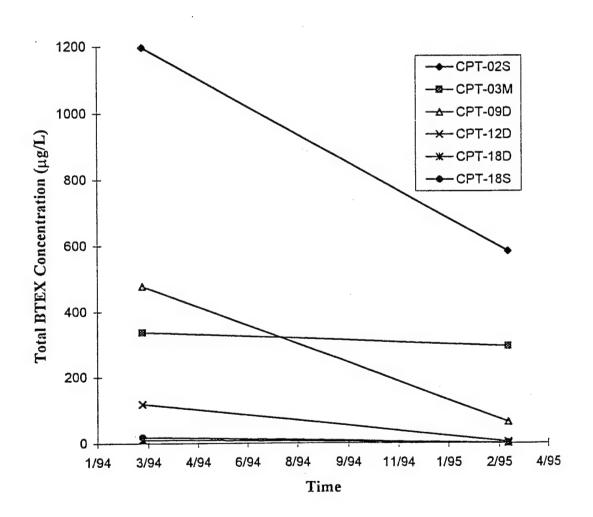
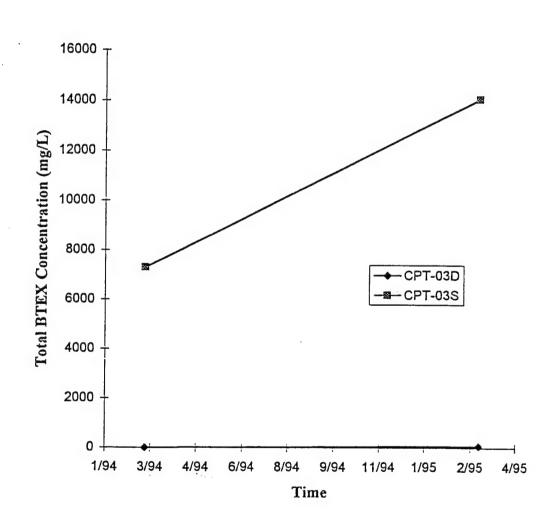


FIGURE 4.5B

# PLOT OF TOTAL BTEX VERSUS TIME FOR SAMPLING LOCATIONS WITH INCREASING BTEX CONCENTRATIONS SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA



# TABLE 4.5

# GROUNDWATER GEOCHEMICAL DATA, MARCH 1994 SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

	Water			Dissolved	Redox	Total	Hydrogen	Ferrous			NO2+NO3		
Sample	Temp.		Conductivity	Oxygen	Potential	Alkalinity	Sulfide	Iron	Chloride	Sulfate	Nitrogen	TOC	Methane
Number	(၁)	Ild	(mp/soum)	(mg/L)	(mV)	(mg/L.)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)
CPT-86-001	24.7	NA V	ΥN	0.4	NA	VN	NA	NA	44.4	4.37	0.13	14	4.99
CPT-86-002D	24.7	7.1	77.1	9.0	-190	330	0.4	0.4	45.7	ND2 °	0.13	8.9	5.953
CPT-86-002S	24.7	6.7	1901	0.2	-156	498	0.2	1.6	42.6	ND2	0.12	16.9	14.953
CPT-86-003D	¥	7.3	721	٧V	-255	315	ı	0.4	41.6	ND2	0.12	5.4	1.63
CPT-86-003M	26.4	7.1	868	0.2	-50	398	0	0.3	40.7	2.52	0.11	10.9	3.164
Cl7T-86-003S	26.4	6.7	1733	0.1	-208	520	0.1	1.2	132	118	14.8	63.1	14.021
CPT-86-0041)	26.1	7.2	457	0.2	-266	212	0.2	9.0	12.4	1.47	0.09	5.6	3.756
CPT-86-004S	26.9	6.9	469	0.3	-286	215	0.5	9.0	12.5	ND2	0.19	9.9	7.661
CPT-86-005S	26.4	7.3	488	1:1	-160	215	0.1	0.1	23.6	98.9	0.17	12	4.858
CPT-86-006S	25.1	7.6	437	0.2	-278	148	1.5	0.3	47.8	7.03	0.13	3.8	6.595
CPT-86-007S	25.3	7.2	577	0.2	-250	254	1.2	-	30.2	2.52	0.12	3.4	6:339
CPT-86-008S	25	7.1	974	0.2	09-	420	0	1.9	44.7	8.51	0.1	10.1	1.742
CPT-86-009D	27.8	7.1	938	0.3	-200	422	9.0	0.2	34.7	15.3	0.11	12	4.236
CPT-86-009S	27.3	7.3	530	0.2	-24	340	0	0.2	14.3	6.64	0.1	10.2	3.797
CPT-86-010S	26	7.3	460	0.1	09-	192	0	0.2	26.6	9.5	0.13	21.3	3.493
CPT-86-011S	25.9	7.2	808	0.1	-35	210	0	0.4	12.7	15.9	0.15	NA	4.244
CPT-86-012D	27.1	7.2	715	0.4	10	329	0.1	0.1	28.1	3.86	0.12	8.1	0.983
CPT-86-012S	27.3	7	564	6.0	30	566	0	0.1	15.2	8.38	0.1	10.5	5.372
CPT-86-013S	25.7	7.3	801	0.1	-230	362	9.0	0.3	35.5	6.94	0.12	7.2	2.043
CPT-86-014D	25.5	7	906	0.3	-240	460	9.0	0.3	34.6	3.68	0.11	12.8	8.793
CPT-86-016DD	26.7	Ϋ́	NA	2.7	NA	NA	0.3	NA	ΝΑ	NA	NA	NA	NA
CPT-86-016S	25.5	7	776	0.1	-190	231	0.1	0.4	37.9	8.23	0.13	9.4	0.781
CPT-86-018D	26.1	7.1	620	0.3	09-	294	0.2	0.3	15	1.85	0.11	5.6	4.56
CPT-86-018DD	27.3	NA	NA	2.3	NA	NA	0	ΑA	NA	ΝA	NA	Ϋ́	NA
CPT-86-018S	26.6	6.9	834	2	25	286	0	0.5	36.6	98	0.12	7.8	ΝΑ
CPT-86-019D	26.6	7.1	744	0.2	-50	328	0.2	0.7	33.7	1.51	0.1	7.3	2.136
CPT-86-019S	AN	7	800	NA	41	335	0	0.1	37.4	8.85	0.11	10.3	0.924
CPT-86-020D	25.7	7.1	842	0.3	4	380	0.1	0.2	52	ND2	0.02	9.8	1.114
CPT-86-020S	25	7	368	1.5	23	148	0	0.3	9.83	25.5	0.1	3.6	1.278
CPT-86-021D	26.4	7.2	716	0.2	-20	304	0.1	0.2	29.8	13.3	0.14	6.1	0.46
CPT-86-021S	26	7.1	610	3.2	20	245	0	0.2	26.6	25.5	0.33	7.6	2.414
CPT-86-022D	25.4	6.9	936	0.1	-287	415	5	0.2	ž	٧V	0.12	11.4	998.0

# TABLE 4.5 (CONCLUDED)

# GROUNDWATER GEOCHEMICAL DATA, MARCH 1994 SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

		Water			Dissolved	Redox	Total	Hydrogen	Ferrous			NO.+NO.		
Sample	Sample	Temp.		Conductivity	Oxygen	Potential	Alkalinity	Sulfide	_	Chloride	Sulfate	Nitrogen	TOC	Methans
Location .	Number	(၃)	Hq	(µmhos/cm)	(mg/L)	(m)	(me/L)	(Dom)	( I/om)	( L'out)	(1/010)	( 1/2 m)		A L
							/ 3	( - G )	( Q	(7,9,)	(111B/12)	(mg/L)	(mg/L)	(mg/L)
CPT-22S	CPT-86-022S	25.5	8.9	1271	0.3	.153	450		1.3	7 7 7 7	961	200		0.00
CPT-23D	CPT-86-023D	267	7.1	770		691	333		7 0	0.00	170	0.07	IO	3.218
CPF.238	CDT 94 0230	3 7 5		100	1.0	2	332	7.0	7.0	30.1	1.49	0.1	8.2	2.282
Chr 245	C1 1-00-1233	50.3	6	/2/	3.5	7.	346	0	0.2	23.4	NI)2	0.12	6.4	1.992
CF 1-24D	CI'I-80-024D	70	7.5	376	0.3	-90	192	0.1	0.1	5.46	3.61	0.1	2.8	0.686
CPT-24S	CPT-86-024S	25.7	7	358	1.7	30	190	0	0.1	6.63	ND2	0.12	2,6	2000
CPT-25D	CPT-86-025D	NA	7.1	892	ΑN	62	371	0	P ICN	547	414	51.0	200	1 657
CPT-25S	CPT-86-025S	25	73	66.1	2.7	13			1277	7.5	0.10	0.12	13.7	1.330
עאר יושט	CD 20 1,00		? .	100	2:/	22	13/	٥	ICIN	28	51.9	0.12	15.7	0.147
(107-1-70)	CI.1-80-070I)	7.07	_	751	0.2	-293	311	e	0.4	44.9	19.8	110	8.3	2 061
CPT-26S	CPT-86-026S	56	7.6	558	2.2	-20	264	C	0.3	- 2	1 22	0.10		200
MW-100	CPT-86-100	25.9	7.2	607	0.5	-2.41	331	0.0	2 -		16.31	21.0	2	3.309
MW-101	CPT-86-101	26.5	7.2	533	0.3	72.	287	8.0	-	21.5	5.01	21.0	18.0	7.871
MW-102	CPT-86-102	25.8	72	523	0.0	201	050	0.0	-	0.12	2.73	0.1.3	6	2.308
MW-103	CDT 96 103	1 36	,		;;;	107	2.30	0.0	1.0	6/1	3.51	0.12	7.6	3.256
201-1414	C11-00-103	47.67	4.	445	0.1	-271	209	_	0.1	12.5	4.69	0.11	26	5 201
1115	CPT-86-PB5	24.9	7.1	743	0.2	-140	291	Ϋ́	ICIN	P IS	4.45	-	VIX.	
PPOL2-1	CPT-86-PPOL2-1	26.3	7.1	747	0.1	-220	305	0.7	90	=	2	133	V <sub>2</sub> 1	4.414
PPOL 2-6	CPT-86-PPOI 2.6	27.2	,	30100	5	020			2		3.6	ICIN	VV	1.33
	0-700 11 00 1 10	- 1	-	20100	7.0	-530	334	8.0	AA	10200	1150	ION	NA	0.034

<sup>\*</sup> TOC = Total organic carbon.
b NA = Not available.

a' NI)2 = <0.5 mg/L. a' NI)1 = <0.05 m/L.

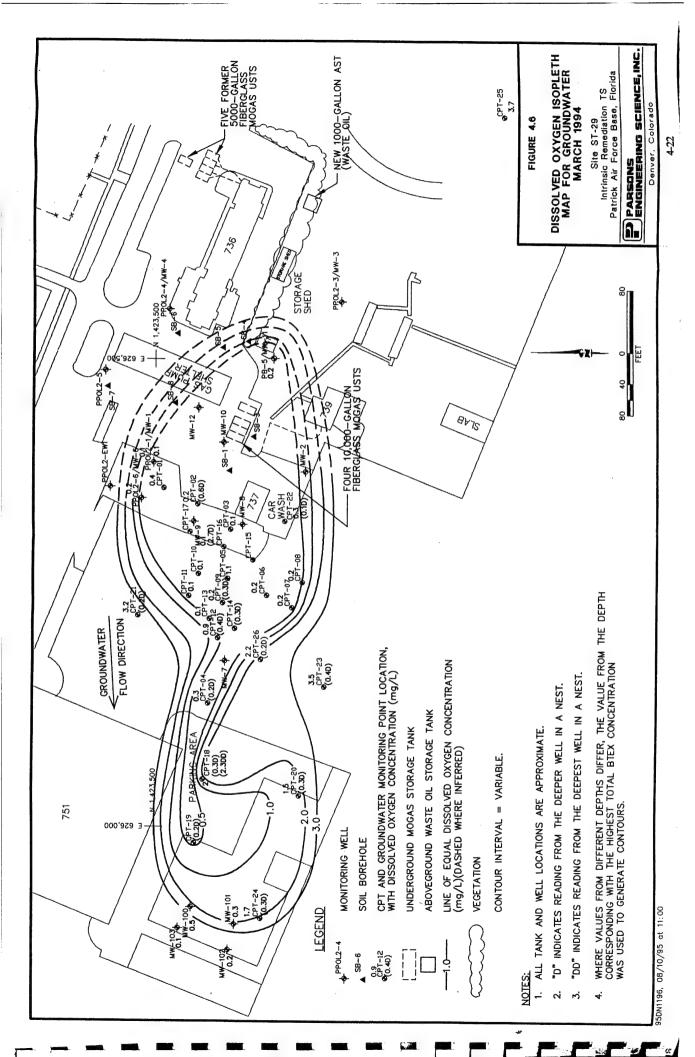
TABLE 4.6

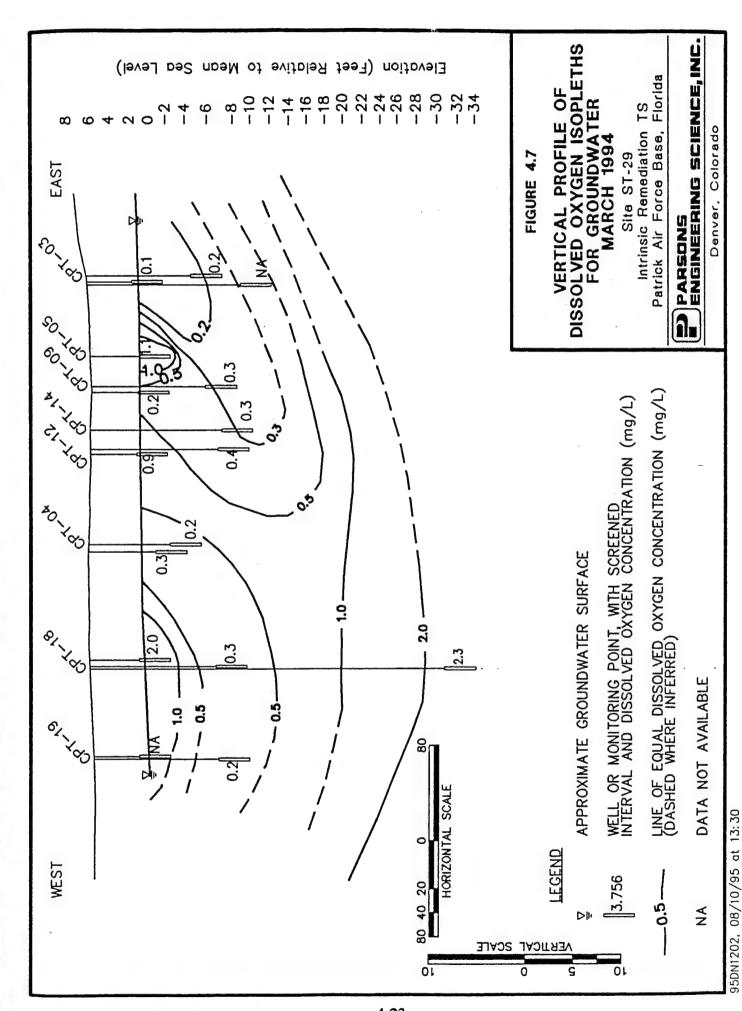
# GROUNDWATER GEOCHEMICAL DATA, MARCH 1995 SITE ST-29 INTRINSIC REMEDIATION TS PATRICK AFB, FLORIDA

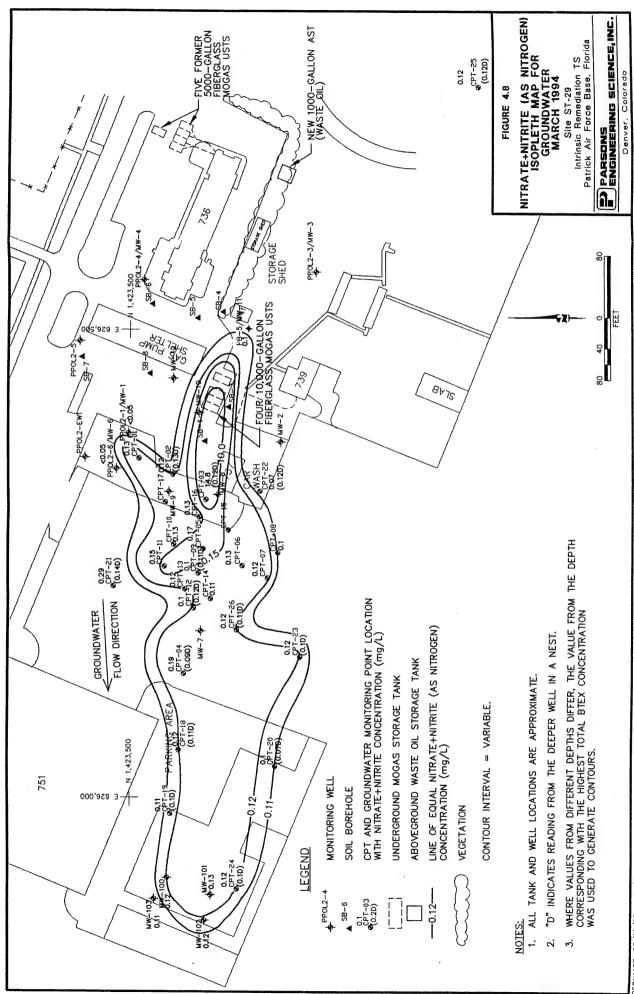
Methane	(mg.L)	12.846	14.15	2.57	٧×	12.437	15.534	9.839	5.822	0.882	12.339	3.756	600.6	5.095	11.63	10.594	0.074	6.116	890.0	3,379	NA	0.749	9.857
Carbon Diavide	(mg/L)	420	416	144	٧×	338	780	284	08	202	142	192	154	06	138	NA	238	86	276	110	132	96	190
Hydrogen Sulfide Carbon Dioxide Methans	(mg/L)	NA.	NA	5	٧٧	NA	NA	5	-	5	-	5	0.7	5	-	NA	NA	2	<0.1	0.7	0.3	5	0.7
Sulfate	(mg/L)	1.13	1.08	15.7	٧٧	19.6	14	2.23	1.52	49.7	86.0	3.31	0.94	2.98	1.17	ΝA	1200	1.08	196	59.6	01	38.8	0.52
Manganese	(mg/L)	<0.1	<0.1	<0.1	٧×	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	٧×	<0.1	<0.1	<0.1	<0.1	<0.1	NA	NA
Ferrous Iron	(mg/L.)	1.1	-:	<0.05	٧X	<0.05	0.1	<0.05	0.1	<0.05	0.3	<0.05	0.3	0.1	0.1	Ϋ́Α	<0.05	0.1	0.1	0.2	0.3	0.1	9.0
Aminonia	(mg/L)	3.83	3.79	1.18	٧V	6.13	18.2	3.85	1.71	2.37	4.04	2.61	1.01	2.53	3.6	Ϋ́	16.9	1.52	16.2	0.35	0.23	0.47	2.94
Nitrate + Nitrite   Ammonia	(mg/L.)	90.0	0.07	80.0	NA	90.0	<0.05	90.0	0.07	0.05	0.07	90.0	0.07	0.07	0.07	NA	<0.05	0.07	<0.05	0.07	0.07	0.07	80.0
Dissolved Oxygen	(mg/L)	0.1	0.1	0.1	NA	<0.1	<0.1	0.1	0.1	<0.1	<0.1	0.1	0.1	<0.1	0.1	NA	0.1	0.1	0.2	0.1	<0.1	<0.1	0.1
Chloride	ات ز	62.7	67.7	46.5	٧V	42.6	129	51.1	9.46	86.9	14.4	51.2	13.2	18.4	15.8	NA	0886	29.6	0806	18.3	15.2	28.2	12.3
Alkalinity	(mg/L CaCO3)	404	407	341	NA	447	920	469	180	359	220	357	202	259	263	NA	307	289	335	276	205	287	177
Redox	(mv)	-231	-225	-348	-347	-343	-357	-293	-279	-340	-306	-342	-569	-325	-289	NA	-288	-316	-171	-287	-253	-307	-239
Conductivity	(µs/cm)	866	\$66	787	786	1019	2010	995	391	946	498	837	440	540	543	NA	26900	652	26500	675	429	674	381
1		6.54	6.53	7.02	7.04	6.77	6.72	6.78	7.38	6.93	7.05	88.9	6.87	7.04	6.85	NA	7.13	6.87	7.19	6.85	7.32	7.18	6.92
Sample	Date	3/23/95	3/23/95	3/23/95	3/23/95	3/23/95	3/23/95	3/25/95	3/25/95	3/25/95	3/25/95	3/26/95	3/26/95	3/25/95	3/23/95	3/23/95	3/31/95	3/24/95	3/31/95	3/24/95	3/23/95	3/25/95	3/25/95
Sample	Number	25	2S DUP	3D	3D DUP	3NI	3.5	9D	95	12D	12S	26D	26S	86-41)	86-45	86-4S DUP	86-16DD	86-MW18D	86-N1W18DD	86-NIW18S	86-NfW100	86-N1W21D	86-NIW21S
Sample	Location	CPT-02S	CPT-02S	CPT-03D			١						CPT-26S	CPT-04D	CPT-04S	CPT-04S	CPT-16DD 86-16DD	$\neg$	اه				CPT-21S

a/ NA = Data not available.

12.00

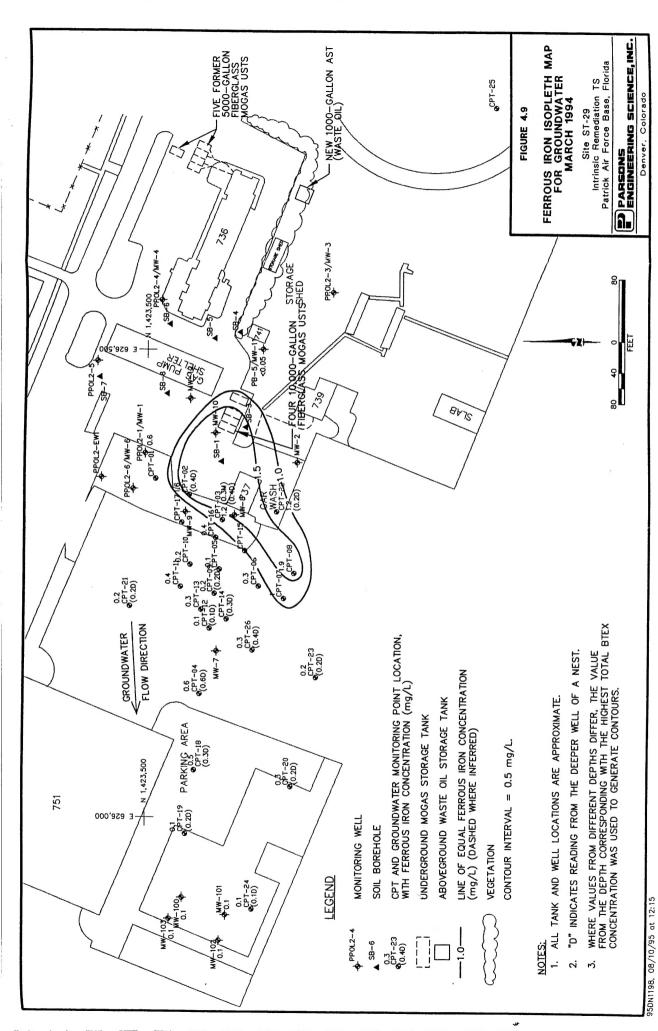


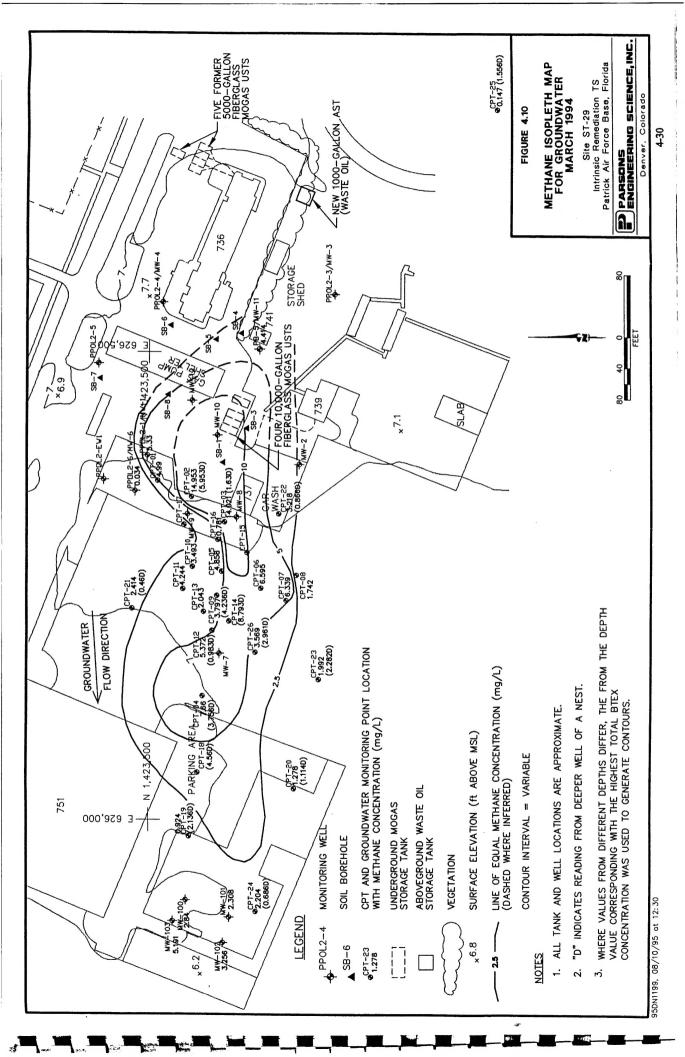


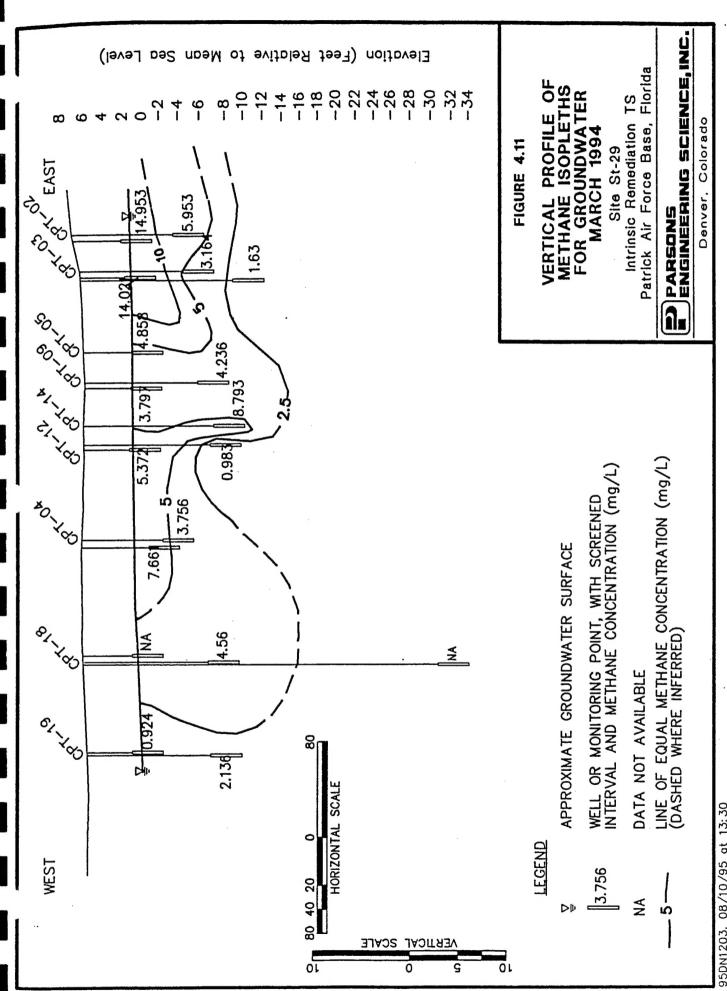


95DN1197, 08/10/95 at 11:30

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95DN1203, 08/10/95 at 13:30

